

Methods of Using Dihydropteridinones

Application Data:

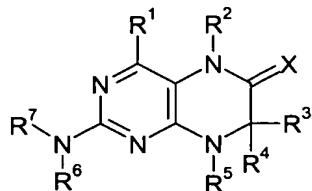
This application is a continuation of US application serial no. 10/226,710 filed

5 08/23/2002 which claims benefit to DE 101 43 272.0 filed 9/4/2001 and US
provisional application no. 60/332681 filed 11/14/2001.

Field of the Invention:

The present invention relates to new dihydropteridinones of general formula (I)

10



wherein the groups X, R¹, R², R³, R⁴, R⁵, R⁶ and R⁷ have the meanings given in

the claims and specification, the isomers thereof, processes for preparing these

15 dihydropteridinones and the use thereof as pharmaceutical compositions.

Background to the invention

Pteridinone derivatives are known from the prior art as active substances with an
20 antiproliferative activity. WO 01/019825 describes the use of pteridinone
derivatives for the treatment of neoplastic and viral diseases. The resistance of
many types of tumours calls for the development of new pharmaceutical
compositions for combating tumours.

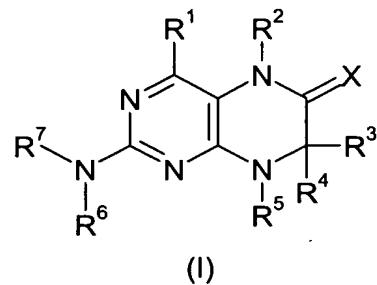
The aim of the present invention is to prepare new compounds with an
25 antiinflammatory and antiproliferative activity.

Detailed description of the invention

Surprisingly it has been found that compounds of general formula (I) wherein the
groups X and R¹ to R⁷ have the meanings given hereinafter act as inhibitors of

specific cell cycle kinases. Thus, the compounds according to the invention may be used for example to treat diseases connected with the activity of specific cell cycle kinases and characterised by excessive or abnormal cell proliferation.

5 The present invention therefore relates to compounds of general formula (I)



wherein

10

R^1 denotes a group selected from among hydrogen, NH_2 , XH , halogen and a C_1 - C_3 -alkyl group optionally substituted by one or more halogen atoms,

15

R^2 denotes a group selected from among hydrogen, CHO , XH , $-X-C_1-C_2$ -alkyl and an optionally substituted C_1 - C_3 -alkyl group,

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R^3 , R^4 which may be identical or different denote a group selected from among optionally substituted C_1 - C_{10} -alkyl, C_2 - C_{10} -alkenyl, C_2 - C_{10} -alkynyl, aryl, heteroaryl, C_3 - C_8 -cycloalkyl, C_3 - C_8 -heterocycloalkyl, $-X$ -aryl, $-X$ -heteroaryl, $-X$ -cycloalkyl, $-X$ -heterocycloalkyl, $-NR^8$ -aryl, $-NR^8$ -heteroaryl, $-NR^8$ -cycloalkyl and $-NR^8$ -heterocycloalkyl, or a group selected from among hydrogen, halogen, $COXR^8$, $CON(R^8)_2$, COR^8 and XR^8 , or R^3 and R^4 together denote a 2- to 5-membered alkyl bridge which may contain 1 to 2 heteroatoms,

25

R⁵ denotes hydrogen or a group selected from among optionally substituted C₁-C₁₀-alkyl, C₂-C₁₀-alkenyl, C₂-C₁₀-alkynyl, aryl, heteroaryl and -C₃-C₆-cycloalkyl , or

5 R³ and R⁵ or R⁴ and R⁵ together denote a saturated or unsaturated C₃-C₄-alkyl bridge which may contain 1 to 2 heteroatoms,

R⁶ denotes optionally substituted aryl or heteroaryl,

R⁷ denotes hydrogen or -CO-X-C₁-C₄-alkyl, and

10 X in each case independently of one another denotes O or S,

R⁸ in each case independently of one another denotes hydrogen or a group selected from among optionally substituted C₁-C₄-alkyl, C₂-C₄-alkenyl, 15 C₂-C₄-alkynyl and phenyl,

optionally in the form of the tautomers, the racemates, the enantiomers, the diastereomers and the mixtures thereof, and optionally the pharmacologically acceptable acid addition salts thereof.

20 Preferred compounds of formula (I) are those wherein X and R⁶ have the meaning indicated, and

R¹ denotes hydrogen,

25 R² denotes a group selected from among a CHO, OH, and CH₃ group,

R³, R⁴ which may be identical or different denote a group selected from among hydrogen, optionally substituted C₁-C₆-alkyl, C₂-C₆-alkenyl, C₂-C₆-alkynyl, C₃-C₇-30 cycloalkyl, or

R³ and R⁴ together denote a C₂-C₅-alkyl bridge ,

R^5 denotes a group selected from among optionally substituted C_1 - C_{10} -alkyl, C_2 - C_{10} -alkenyl, C_2 - C_{10} -alkynyl, C_3 - C_6 -cycloalkyl and C_3 - C_6 -cycloalkenyl, or R^3 and R^5 or R^4 and R^5 together denote a saturated or unsaturated C_3 - C_4 -alkyl bridge which may contain 1 to 2 heteroatoms, and

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R^7 denotes hydrogen,

optionally in the form of the tautomers, the racemates, the enantiomers, the diastereomers and the mixtures thereof, and optionally the pharmacologically

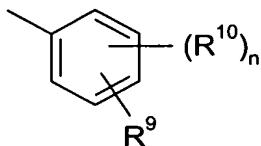
10 acceptable acid addition salts thereof.

Particularly preferred compounds of formula (I) are those wherein

R^1 - R^5 , R^7 , R^8 and X have the meaning indicated, and

R^6 denotes a group of general formula

15



wherein

n denotes 1, 2, 3 or 4,

20

R^9 denotes a group selected from among optionally substituted C_1 - C_6 -alkyl, C_2 - C_6 -alkenyl, C_2 - C_6 -alkynyl, $-CONH-C_1-C_{10}$ -alkylene, $-O$ -aryl, $-O$ -heteroaryl, $-O$ -cycloalkyl, $-O$ -heterocycloalkyl, aryl, heteroaryl, cycloalkyl and heterocycloalkyl or a group selected from among $-O-C_1-C_6$ -alkyl- Q^1 , $-CONR^8-C_1-C_{10}$ -alkyl- Q^1 , $-CONR^8-C_2-C_{10}$ -alkenyl- Q^1 , $-CONR^8-Q^2$, halogen, OH, $-SO_2R^8$, $-SO_2N(R^8)_2$, $-COR^8$, $-COOR^8$, $-N(R^8)_2$, $-NHCOR^8$, $CONR^8OC_1-C_{10}$ alkyl- Q^1 and $CONR^8OQ^2$,

25

Q^1 denotes hydrogen, $-NHCOR^8$, or a group selected from among an optionally substituted $-NH$ -aryl, $-NH$ -heteroaryl, aryl, heteroaryl, C_3 - C_8 -cycloalkyl- and heterocycloalkyl group,

Q^2 denotes hydrogen or a group selected from among an optionally substituted aryl, heteroaryl, C_3 - C_8 -heterocycloalkyl, C_3 - C_8 -cycloalkyl- and C_1 - C_4 -alkyl- C_3 - C_8 -cycloalkyl group,

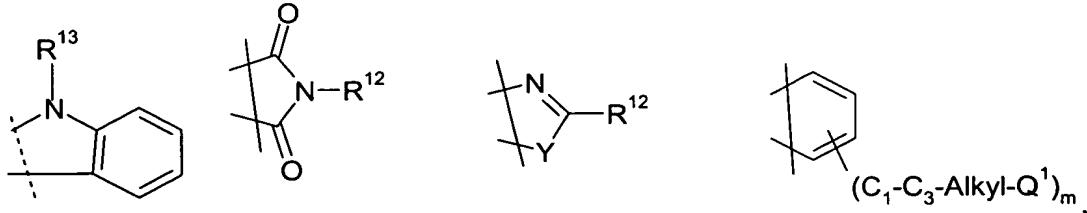
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R^{10} which may be identical or different denotes a group selected from among optionally substituted C_1 - C_6 -alkyl, C_2 - C_6 -alkenyl and C_2 - C_6 -alkynyl, $-O-C_1-C_6$ -alkyl, $-O-C_2-C_6$ -alkenyl, $-O-C_2-C_6$ -alkynyl, C_3 - C_6 -heterocycloalkyl and C_3 - C_6 -cycloalkyl, or a group selected from among hydrogen, $-CONH_2$, $-COOR^8$, $-OCON(R^8)_2$,

10 $-N(R^8)_2$, $-NHCOR^8$, $-NHCON(R^8)_2$, $-NO_2$ and halogen,

or

adjacent groups R^9 and R^{10} together denote a bridge of general formula



Y denotes O, S or NR^{11} ,

15 m denotes 0, 1 or 2

R^{11} denotes hydrogen or C_1 - C_2 -alkyl, and

R^{12} denotes hydrogen or a group selected from among optionally substituted

20 phenyl, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, $-C_1-C_3$ -alkyl-phenyl, $-C_1-C_3$ -alkyl-pyridyl, $-C_1-C_3$ -alkyl-pyrazinyl, $-C_1-C_3$ -alkyl-pyrimidinyl and $-C_1-C_3$ -alkyl-pyridazinyl,

R^{13} denotes C_1 - C_6 -alkyl,

25 optionally in the form of the tautomers, the racemates, the enantiomers, the diastereomers and the mixtures thereof, and optionally the pharmacologically acceptable acid addition salts thereof.

30 Particularly preferred are compounds of formula (I) wherein

R^3 - R^6 , R^8 and X have the meaning indicated, and

R^1 denotes hydrogen,

R^2 denotes CH_3 , and

R^7 denotes hydrogen,

5 optionally in the form of the tautomers, the racemates, the enantiomers, the diastereomers and the mixtures thereof, and optionally the pharmacologically acceptable acid addition salts thereof.

The invention further relates to compounds of formula (I), wherein X and R^1 - R^7

10 have the meanings indicated, for use as pharmaceutical compositions.

Of particular importance according to the invention are compounds of formula (I), wherein X and R^1 - R^7 have the meaning indicated, for use as pharmaceutical compositions with an antiproliferative activity.

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The invention also relates to the use of a compound of formula (I), wherein X and R^1 - R^7 have the meaning indicated, for preparing a pharmaceutical composition for the treatment and/or prevention of cancer, infections, inflammatory and autoimmune diseases.

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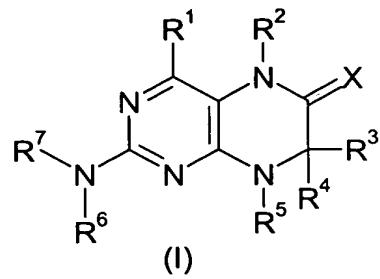
The invention also relates to a method of treating and/or preventing cancer, infections, inflammatory and autoimmune diseases, characterised in that a patient is given an effective amount of a compound of formula (I), wherein X and R^1 - R^7 have the meanings indicated.

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The invention also relates to pharmaceutical preparations, containing as active substance one or more compounds of general formula (I), wherein X and R^1 - R^7 have the meaning indicated, or the physiologically acceptable salts thereof, optionally combined with conventional excipients and/or carriers.

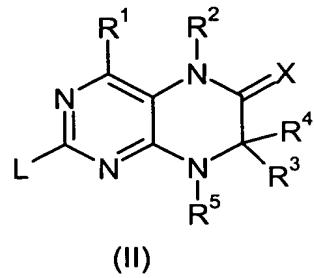
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The invention also relates to a process for preparing a compound of general formula (I),



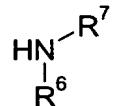
wherein

5 R¹-R⁷ and X are as hereinbefore defined,
characterised in that a compound of general formula (II)



wherein

R¹-R⁵ and X are as hereinbefore defined and L is a leaving group,
10 is reacted with an optionally substituted compound of general formula (III)

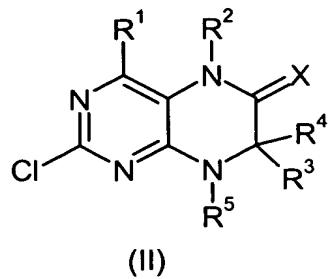


(III)

wherein

15 R⁶ and R⁷ are as hereinbefore defined.

The invention also relates to a compound of formula (II),

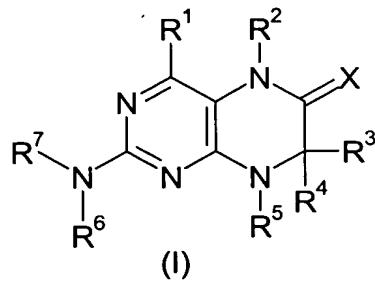


wherein

R^1 - R^5 and X are as hereinbefore defined. Compounds of formula (II) are important intermediate products for preparing the compounds of formula (I) according to the invention.

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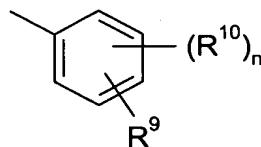
The invention also relates to a process for preparing a compound of general formula (I),



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wherein

R^6 denotes a group of general formula,

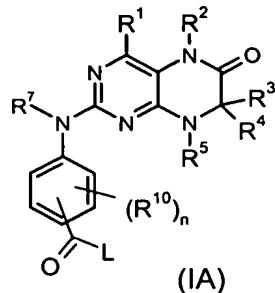


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R^9 denotes an optionally substituted group -CONH-C₁-C₁₀-alkylene or a group selected from among -CONR⁸-C₁-C₁₀-alkyl-Q¹, -CONR⁸-C₂-C₁₀-alkenyl-Q¹, -CONR⁸-Q² and -COOR⁸, and

20 R^1 - R^5 , R^7 , R^{10} , n and X are as hereinbefore defined,

characterised in that a compound of general formula (IA)



wherein

5 R¹ to R⁵, R⁷ and R¹⁰ are as hereinbefore defined, and
L denotes a leaving group,

is reacted with a primary or secondary amine to form the corresponding amide or
is reacted with an alcohol to form the corresponding ester.

10

The term alkyl groups, including alkyl groups which are a part of other groups, denotes branched and unbranched alkyl groups with 1 to 10 carbon atoms, preferably 1 – 6, most preferably 1-4 carbon atoms, such as, for example: methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, nonyl and decyl. Unless otherwise stated, the abovementioned terms propyl, butyl, pentyl, hexyl, heptyl, octyl, nonyl and decyl include all the possible isomeric forms. For example, the term propyl includes the two isomeric groups n-propyl and iso-propyl, the term butyl includes n-butyl, iso-butyl, sec. butyl and tert.-butyl, the term pentyl includes iso-pentyl, neopentyl, etc.

20

In the abovementioned alkyl groups one or more hydrogen atoms may optionally be replaced by other groups. For example these alkyl groups may be substituted by the halogen atoms fluorine, chlorine, bromine or iodine. The substituents fluorine and chlorine are preferred. The substituent chlorine is particularly preferred. All the hydrogen atoms of the alkyl group may optionally also be replaced.

Similarly, in the abovementioned alkyl groups, unless otherwise stated, one or more hydrogen atoms may optionally be replaced for example by an optionally

substituted group selected from among CN, OCOCH₃, aryl, preferably phenyl, heteroaryl, preferably thienyl, thiazolyl, imidazolyl, pyridyl, pyrimidyl or pyrazinyl, saturated or unsaturated heterocycloalkyl, preferably pyrazolyl, pyrrolidinyl, piperidinyl, piperazinyl or tetrahydro-oxazinyl, an amine group, preferably

5 methylamine, benzylamine, phenylamine or heteroarylamine, saturated or unsaturated bicyclic ring systems, preferably benzimidazolyl and cycloalkyl, preferably cyclohexyl or cyclopropyl.

The term alkyl bridge, unless otherwise stated, denotes branched and unbranched

10 alkyl groups with 2 to 5 carbon atoms, for example propylene, isopropylene, n-butylene, iso-butyl, sec. butyl and tert.-butyl etc. bridges. Propylene and butylene bridges are particularly preferred. In the alkyl bridges mentioned 1 to 2 C-atoms may optionally be replaced by one or more heteroatoms selected from among oxygen, nitrogen or sulphur.

15

The term alkenyl groups (including those which are a part of other groups) denotes branched and unbranched alkylene groups with 2 to 10 carbon atoms, preferably 2 - 6 carbon atoms, most preferably 2 - 3 carbon atoms, provided that they have at least one double bond. Examples include: ethenyl, propenyl, butenyl, pentenyl

20 etc. Unless otherwise stated, the abovementioned terms propenyl, butenyl, etc also include all the possible isomeric forms. For example, the term butylene includes n-butenyl, 1-methylpropenyl, 2-methylpropenyl, 1,1-dimethylethenyl, 1,2-dimethylethenyl etc.

25 In the abovementioned alkenyl groups, unless otherwise stated, one or more hydrogen atoms may optionally be replaced by other groups. For example, these alkyl groups may be substituted by the halogen atoms fluorine, chlorine, bromine or iodine. The substituents fluorine and chlorine are preferred. The substituent chlorine is particularly preferred. All the hydrogen atoms of the alkenyl group may optionally also be replaced.

30

The term alkynyl groups (including those which are a part of other groups) denotes branched and unbranched alkynyl groups with 2 to 10 carbon atoms, provided that

they have at least one triple bond, for example ethynyl, propargyl, butynyl, pentynyl, hexynyl etc., preferably ethynyl or propynyl.

In the abovementioned alkynyl groups, unless otherwise stated, one or more

5 hydrogen atoms may optionally be replaced by other groups. For example, these alkyl groups may be substituted by the halogen atoms fluorine, chlorine, bromine or iodine. The substituents fluorine and chlorine are preferred. The substituent chlorine is particularly preferred. All the hydrogen atoms of the alkynyl group may optionally also be replaced.

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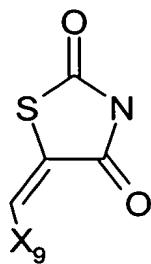
The term aryl denotes an aromatic ring system with 6 to 14 carbon atoms, preferably 6 or 10 carbon atoms, preferably phenyl, which, unless otherwise stated, may carry one or more of the following substituents, for example: OH, NO₂,

15 CN, -OCHF₂, -OCF₃, -NH₂, halogen, for example fluorine, chlorine, bromine or iodine, preferably fluorine or chlorine, C₁-C₁₀-alkyl, preferably C₁-C₅-alkyl, preferably C₁-C₃-alkyl, most preferably methyl or ethyl, -O-C₁-C₃-alkyl, preferably -O-methyl or -O-ethyl, -N-methyl -tetrahydro-oxazinyl, -COOH, -COO-C₁-C₄-alkyl, preferably -COOCH₂CH₃, -COO-C(CH₃)₃ or -COOCH₃, -CONH₂,

-CONH-C₁-C₁₀-alkyl, while this alkyl may optionally be further substituted,

20 optionally substituted -CONH-C₃-C₆-cycloalkyl, preferably optionally substituted -CONH-cyclopentyl, optionally substituted -CONH-heterocycloalkyl, preferably piperidinyl, pyrrolidinyl or piperazinyl, optionally substituted -CONH-heteroaryl, preferably optionally substituted -CONH-pyridyl, optionally substituted -CONH-aryl, preferably optionally substituted -CONH-phenyl, -CONMeC₁-C₃-alkyl, while 25 this alkyl may optionally be further substituted, preferably -CONMeCH₂-pyridyl, benzimidazole or

a group of formula

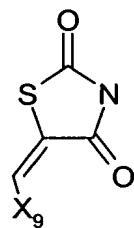


5 Examples of 5-10-membered mono- or bicyclic heteroaryl rings wherein up to three C-atoms may be replaced by one or more heteroatoms selected from among oxygen, nitrogen or sulphur include furan, thiophene, pyrrole, pyrazole, imidazole, triazole, tetrazole, pyridine, pyridazine, pyrimidine, pyrazine, triazine, oxazole, isoxazole, thiazole, thiadiazole and oxadiazole, while each of the abovementioned

10 heterocycles may optionally also be annellated onto a benzene ring, preferably benzimidazole, and unless otherwise stated these heterocycles may for example carry one or more of the following substituents: OH, NO₂, CN, -OCHF₂, -OCF₃, -NH₂, halogen, preferably fluorine or chlorine, C₁-C₁₀-alkyl, preferably C₁-C₅-alkyl, preferably C₁-C₃-alkyl, most preferably methyl or ethyl, -O-C₁-C₃-alkyl, preferably -

15 O-methyl or -O-ethyl, -methyl-N-tetrahydro-oxazinyl, -COOH, -COO-C₁-C₄-alkyl, preferably -COO-C(CH₃)₃ or -COOCH₃, -CONH₂, optionally substituted phenyl, optionally substituted heteroaryl, preferably optionally substituted pyridyl or pyrazinyl, -CONH-C₁-C₁₀-alkyl, while this alkyl may itself optionally be substituted, optionally substituted -CONH-C₃-C₆-cycloalkyl, preferably optionally substituted -

20 CONH-cyclopentyl, optionally substituted -CONH-heteroaryl, preferably optionally substituted -CONH-pyridyl, optionally substituted -CONH-aryl, preferably optionally substituted -CONH-phenyl, -CONMeC₁-C₃-alkyl, while this alkyl may itself optionally be substituted, preferably -CONMeCH₂-pyridyl, benzimidazole or a group of formula



The term cycloalkyl groups denotes, for example, saturated or unsaturated cycloalkyl groups with 3 - 8 carbon atoms, for example cyclopropyl, cyclobutyl,

5 cyclopentyl, cyclopentenyl, cyclohexyl, cyclohexenyl, cycloheptyl or cyclooctyl, preferably cyclopropyl, cyclopentyl or cyclohexyl, while each of the abovementioned cycloalkyl groups may optionally also carry one or more substituents, preferably =O, or may be annellated to a benzene ring.

10 "=O" denotes an oxygen atom linked via a double bond.

The term heterocycloalkyl groups, unless otherwise described in the definitions, may denote 5-, 6- or 7-membered, saturated or unsaturated heterocycles, which may contain nitrogen, oxygen or sulphur as heteroatoms, for example

15 tetrahydrofuran, tetrahydrofuranon, γ -butyrolactone, α -pyran, γ -pyran, dioxolane, tetrahydropyran, dioxane, dihydrothiophene, thiolan, dithiolan, pyrrolidine, pyrrolidine, pyrazoline, pyrazolidine, imidazoline, imidazolidine, tetrazole, piperidine, pyridazine, pyrimidine, pyrazine, piperazine, triazine, tetrazine, morpholine, thiomorpholine, diazepan, oxazine, tetrahydro-oxazinyl, isothiazole

20 and pyrazolidine, preferably pyrazolyl, pyrrolidinyl, piperidinyl, piperazinyl or tetrahydro-oxazinyl, while the heterocycle may optionally be substituted.

Generally, the term halogen denotes fluorine, chlorine, bromine or iodine.

25 The leaving group L denotes either identical or different leaving groups such as for example chlorine, bromine, iodine, methanesulphonyl, trifluoromethanesulphonyl or p-toluenesulphonyl, preferably chlorine.

30 The compounds according to the invention may be present in the form of the individual optical isomers, mixtures of the individual enantiomers, diastereomers or

racemates, in the form of the tautomers and also in the form of the free bases or the corresponding acid addition salts with pharmacologically acceptable acids - such as for example acid addition salts with hydrohalic acids, for example hydrochloric or hydrobromic acid, or organic acids, such as for example oxalic,

5 fumaric, diglycolic or methanesulphonic acid.

The substituent R¹ may denote a group selected from among hydrogen, NH₂, XH, preferably OH, halogen, preferably fluorine or chlorine and a C₁-C₃-alkyl group optionally substituted by one or more, preferably one, two or three halogen atoms,

10 preferably fluorine or chlorine, preferably methyl or ethyl. Most preferably, the substituent R¹ is hydrogen .

The substituent R² may denote a group selected from among hydrogen, CHO, XH, preferably OH, -X-C₁-C₂-alkyl, preferably -O-CH₃ or -O-CH₂CH₃, and an

15 optionally substituted C₁-C₃-alkyl group, while the alkyl group preferably consists of 1 to 2 carbon atoms, particularly preferably a carbon atom and may optionally be substituted, preferably by halogen atoms, most preferably by fluorine atoms. In particular, the substituent R² denotes methyl.

20 The substituents R³ and R⁴ may be identical or different and may represent a group selected from among optionally substituted C₁-C₁₀-alkyl, preferably C₁-C₆-alkyl, preferably C₁-C₄-alkyl, most preferably methyl, ethyl or propyl, particularly preferably methyl or ethyl, C₂-C₁₀-alkenyl, preferably ethenyl or propenyl, preferably ethenyl, C₂-C₁₀-alkynyl, preferably ethynyl or propynyl, aryl, preferably 25 optionally substituted phenyl, heteroaryl, C₃-C₈-cycloalkyl, preferably cyclopropyl and cyclobutyl, C₃-C₈-heterocycloalkyl, -X-aryl, -X-heteroaryl, -X-cycloalkyl, -X-heterocycloalkyl, -NR⁸-aryl, -NR⁸-heteroaryl, -NR⁸-cycloalkyl and -NR⁸-heterocycloalkyl, or

30 a group selected from among hydrogen, halogen, COXR⁸, CON(R⁸)₂, COR⁸ and XR⁸, preferably hydrogen, or

the groups R³ and R⁴ may together denote a 2- to 5-membered alkyl bridge, preferably an ethylene, propylene or butylene bridge , while the propylene or

butylene bridge may contain 1 to 2 heteroatoms, preferably oxygen, nitrogen or sulphur, most preferably an ethylene bridge.

Most preferably, the substituent R³ denotes methyl or ethyl. The substituent R⁴ most preferably denotes hydrogen or methyl. Particularly preferred are compounds wherein R³ and R⁴ represent methyl.

All the groups mentioned in the definition of R³ and R⁴ may optionally be substituted.

The group R⁵ may contain hydrogen or a group selected from among optionally substituted C₁-C₁₀-alkyl, for example C₁-C₆-alkyl-aryl or C₁-C₆-alkyl-heteroaryl, preferably C₁-C₆-alkyl, most preferably C₁-C₅-alkyl, particularly preferably propyl, butyl, pentyl, hexyl, -CH₂-cyclohexyl, (CH₂)₁₋₂cyclopropyl or (CH₂)₄-OCOCH₃, C₂-C₁₀-alkenyl, preferably propenyl, butenyl, pentenyl or hexenyl, preferably propenyl or hexenyl, C₂-C₁₀-alkynyl, preferably propynyl, butynyl or pentynyl, preferably propynyl, aryl, preferably phenyl, heteroaryl, -C₃-C₆-cycloalkyl, preferably cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl and -C₃-C₆-cycloalkenyl, preferably cyclohexenyl or cyclopentenyl, or the substituents R³ and R⁵ or R⁴ and R⁵ together denote a saturated or unsaturated C₃-C₄-alkyl bridge which may contain 1 to 2 heteroatoms, preferably oxygen, sulphur or nitrogen.

All the groups mentioned in the definition of R⁵ may optionally be substituted.

The substituent R⁶ may denote optionally substituted aryl, or heteroaryl, preferably aryl, preferably phenyl.

Most preferably, the substituent R⁶ denotes a phenyl group, which may be substituted by one of the groups R⁹ and R¹⁰ described hereinafter, while the phenyl ring may carry one of the groups R⁹, preferably in the *para* position, and one, two, three or four, preferably one or two, of the groups R¹⁰, preferably in the *ortho* or *meta* position.

30

The substituent R⁷ may denote hydrogen or -CO-X-C₁-C₄-alkyl, preferably hydrogen.

X denotes, in each case independently of one another, O or S, preferably O.

The groups R⁸ mentioned in the definitions of the substituents R³ and R⁴

5 represent, independently of one another in each case, hydrogen or a group selected from among optionally substituted C₁-C₄-alkyl, C₂-C₄-alkenyl, C₂-C₄-alkynyl and phenyl, preferably hydrogen or C₁-C₂-alkyl.

The substituent R⁹ may represent a group selected from among optionally

10 substituted C₁-C₆-alkyl, preferably C₁-C₄-alkyl, preferably methyl, ethyl or propyl, most preferably methyl, C₂-C₆-alkenyl, C₂-C₆-alkynyl, -CONH-C₁-C₁₀-alkylene, preferably -CONH-C₁-C₃-alkylene, preferably -CONH-C₁-C₂-alkylene, -O-aryl, preferably O-C₆-C₁₀-aryl, most preferably O-phenyl, -O-heteroaryl, -O-cycloalkyl, preferably O-C₃-C₆-cycloalkyl, most preferably O-cyclopropyl, -O-heterocycloalkyl,

15 aryl, preferably C₆-C₁₀-aryl, most preferably phenyl, heteroaryl, cycloalkyl, preferably C₃-C₆-cycloalkyl, most preferably cyclopropyl, and heterocycloalkyl, or a group selected from among -O-C₁-C₆-alkyl-Q¹, -CONR⁸-C₁-C₁₀-alkyl-Q¹, -CONR⁸-C₁-C₁₀-alkenyl-Q¹, -CONR⁸-Q², halogen, for example fluorine, chlorine, bromine or iodine, OH, -SO₂R⁸, -SO₂N(R⁸)₂, -COR⁸, -COOR⁸, -N(R⁸)₂, -NHCOR⁸,

20 CONR⁸OC₁-C₁₀-alkylQ¹ and CONR⁸OQ², where Q¹ and Q² are as hereinbefore defined.

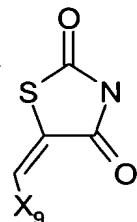
Preferably, R⁹ denotes one of the following groups -CONH-C₁-C₁₀-alkyl, preferably -CONH-C₁-C₃-alkyl, most preferably -CONH-C₁-C₂-alkyl, while this alkyl may itself optionally be substituted, by CN, optionally substituted aryl,

25 preferably optionally substituted phenyl, heteroaryl, preferably thienyl, thiazolyl, imidazolyl, pyridyl, pyrimidyl or pyrazinyl, saturated or unsaturated heterocycloalkyl, preferably pyrazolyl, pyrrolidinyl, piperidinyl, piperazinyl or tetrahydro-oxazinyl, an amine group, preferably methylamine, benzylamine, phenylamine or heteroarylamine, saturated or unsaturated bicyclic ring systems,

30 preferably benzimidazolyl and cycloalkyl, preferably cyclohexyl.

Moreover R⁹ preferably denotes -CONH-heteroaryl, preferably -CONH-pyridyl, -CONH-C₃-C₁₀-cycloalkyl, preferably -CONH-cyclopropyl -CONH-cyclobutyl or

-CONH-cyclopentyl, most preferably -CONH-cyclopropyl; -CONH-C₃-C₁₀-heterocycloalkyl, -CONH-C₆-C₁₀-aryl, preferably -CONH-phenyl, COO-C₁-C₃-alkyl, most preferably COOCH₃, COOH, halogen, preferably F or chlorine, OH or a group of formula



5

All the groups mentioned in the definition of R⁹ may optionally be substituted, preferably by one or more of the groups selected from among OH, OCH₃, Cl, F, CH₃, COOH, CONHCH₂Ph and CONHCH₂-pyrazinyl-CH₃.

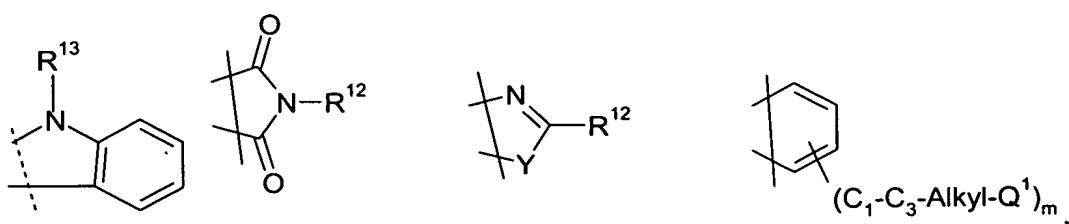
10

The substituent R¹⁰ may be identical or different in each case and may denote a group selected from among optionally substituted C₁-C₆-alkyl, preferably C₁-C₃-alkyl, C₂-C₆-alkenyl, preferably C₂-C₃-alkenyl and C₂-C₆-alkynyl, preferably C₂-C₃-alkynyl, -O-C₁-C₆-alkyl, preferably -O-C₁-C₃-alkyl, -O-C₂-C₆-alkenyl, -O-C₂-C₆-alkynyl, C₃-C₆-heterocycloalkyl and C₃-C₆-cycloalkyl, or a group selected from among hydrogen, -CONH₂, -COOR⁸, -OCON(R⁸)₂, -N(R⁸)₂, -NHCOR⁸, -NHCON(R⁸)₂, -NO₂ and halogen, for example fluorine, chlorine, bromine or iodine.

15

Preferably, the substituent R¹⁰ denotes hydrogen, methyl, methoxy, fluorine or chlorine, most preferably hydrogen or methoxy, particularly preferably methoxy.

Adjacent groups R⁹ and R¹⁰ may together denote a bridge of general formula



wherein

Y denotes O, S or NR¹¹, preferably NR¹¹,

m denotes 0, 1 or 2, preferably 1,

5 R¹¹ denotes hydrogen or C₁-C₂-alkyl, preferably hydrogen or methyl, most preferably hydrogen,

R¹² denotes hydrogen or a group selected from among optionally substituted phenyl, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, -C₁-C₃-alkyl-phenyl, -C₁-C₃-alkyl-

10 pyridyl, -C₁-C₃-alkyl-pyrazinyl, -C₁-C₃-alkyl-pyrimidinyl and -C₁-C₃-alkyl-pyridazinyl, preferably phenyl, pyridyl and pyrazinyl, and

R¹³ denotes C₁-C₆-alkyl, preferably methyl or ethyl.

The compounds according to the invention may be prepared by synthesis methods

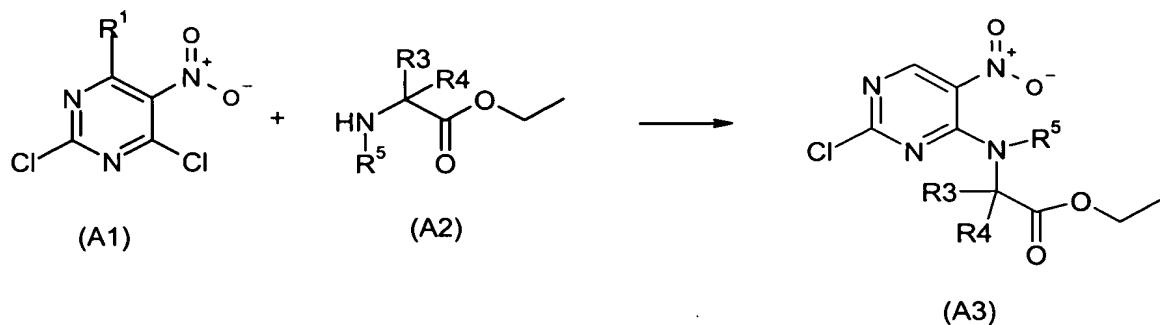
15 A and B described hereinafter, while the substituents of general formulae (A1) to (A6) have the meanings given hereinbefore. These methods are to be understood as illustrations of the invention without restricting it to their subject matter.

Method A

Step 1A

A compound of formula (A1) is reacted with a compound of formula (A2) to obtain a compound of formula (A3) (Diagram 1A). This reaction may be carried out according to WO 0043369 or WO 0043372. Compound (A1) is commercially obtainable, for example, from City Chemical LLC, 139 Allings Crossing Road,

25 West Haven, CT, 06516, USA. Compound (A2) may be prepared by procedures known from the literature: (a) F. Effenberger, U. Burkhardt, J. Willfahrt *Liebigs Ann. Chem.* **1986**, 314-333; b) T. Fukuyama, C.-K. Jow, M. Cheung, *Tetrahedron Lett.* **1995**, 36, 6373-6374; c) R. K. Olsen, *J. Org. Chem.* **1970**, 35, 1912-1915; d) F.E. Dutton, B.H. Byung *Tetrahedron Lett.* **1998**, 30, 5313-5316; e) J. M. Ranajuhi, M. 30 M. Joullie *Synth. Commun.* **1996**, 26, 1379-1384.).

Diagram 1A

5 In Step 1A, 1 equivalent of the compound (A1) and 1 to 1.5 equivalents, preferably 1.1 equivalents of a base, preferably potassium carbonate, potassium hydrogen carbonate, sodium carbonate or sodium hydrogen carbonate, calcium carbonate, most preferably potassium carbonate, are stirred in a diluent, for example acetone, aqueous acetone, tetrahydrofuran, diethylether or dioxane, preferably acetone or diethylether, most preferably acetone.

10 At a temperature of 0 to 15 °C, preferably 5 to 10 °C, 1 equivalent of an amino acid of formula (A2), dissolved in an organic solvent, for example acetone, tetrahydrofuran, diethylether or dioxane, preferably acetone, is added dropwise. The reaction mixture is heated to a temperature of 18°C to 30 °C, preferably about

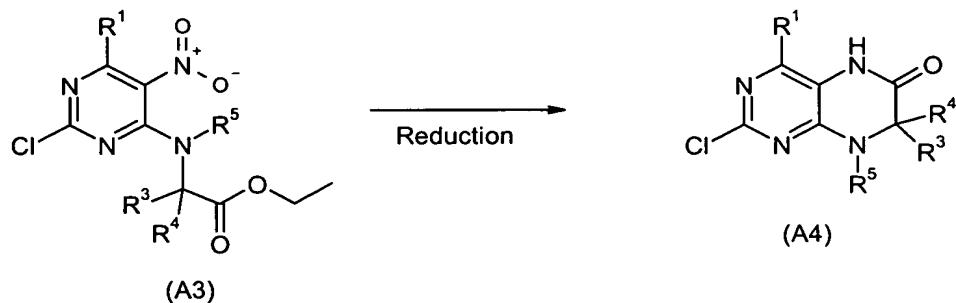
15 22°C, with stirring and then stirred for a further 10 to 24 hours, preferably about 12 hours. Then the diluent is distilled off, the residue is combined with water and the mixture is extracted two to three times with an organic solvent, such as diethylether or ethyl acetate, preferably ethyl acetate. The combined organic extracts are dried and the solvent is distilled off. The residue (compound A3) may

20 be used in Step 2 without any prior purification.

Step 2A

The compound obtained in Step 1A (A3) is reduced at the nitro group and cyclised to form the compound of formula (A4) (Diagram 2A).

5

Diagram 2A

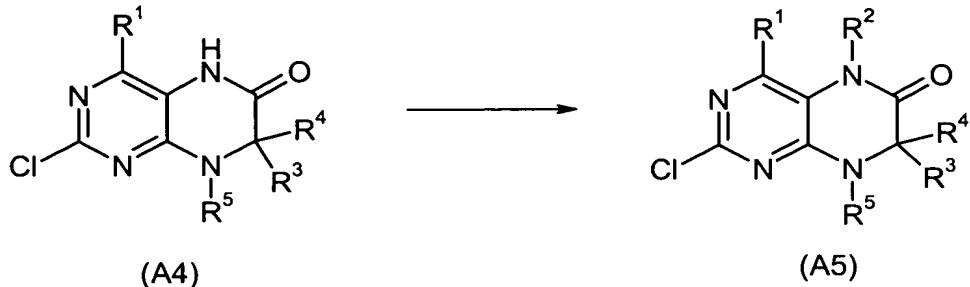
In Step 2A , 1 equivalent of the nitro compound (A3) is dissolved in an acid,

- 10 preferably glacial acetic acid, formic acid or hydrochloric acid, preferably glacial acetic acid, and heated to 50 to 70 °C, preferably about 60 °C. Then a reducing agent, for example zinc, tin or iron, preferably iron filings, is added to complete the exothermic reaction and the mixture is stirred for 0.2 to 2 hours, preferably 0.5 hours, at 100 to 125 °C, preferably at about 117 °C. After cooling to ambient
- 15 temperature the iron salt is filtered off and the solvent is distilled off. The residue is taken up in a solvent or mixture of solvents, for example ethyl acetate or dichloromethane/ methanol 9/1 and semisaturated NaCl solution, and filtered through kieselgur, for example. The organic phase is dried and evaporated down. The residue (compound (A4)) may be purified by chromatography or by
- 20 crystallisation or used as the crude product in Step 3A of the synthesis.

Step 3A

The compound obtained in Step 2A (A4) may be reacted by electrophilic substitution as shown in Diagram 3A to obtain the compound of formula (A5).

5 Diagram 3A

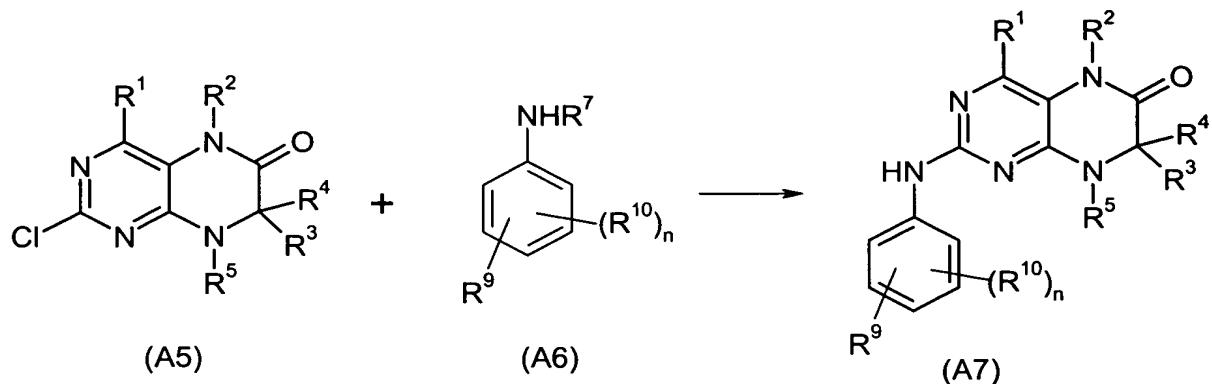


In Step 3A 1 equivalent of the amide of formula (A4) is dissolved in an organic solvent, for example dimethylformamide or dimethylacetamide, preferably dimethylacetamide, and cooled to about -5 to 5 °C, preferably 0 °C.

10 Then 0.9 to 1.3 equivalents of sodium hydride and 0.9 to 1.3 equivalents of alkyl
halide, for example methyl iodide, are added. The reaction mixture is stirred for 0.1
– 3 hours, preferably about 1 hour, at about 0 to 10 °C, preferably at about 5 °C,
and may optionally be left to stand for a further 12 hours at this temperature. The
reaction mixture is evaporated down and extracted with water and an organic
15 solvent, preferably dichloromethane or ethyl acetate. The organic phases are
evaporated down. The residue (compound (A5)) may be purified by
chromatography, preferably over silica gel.

20 Step 4A

The amination of the compound (A5) obtained in Step 3A to yield the compound of formula (A7) (Diagram 4A) may be carried out using the methods known from the literature of variants 4.1 A (a) M.P.V. Boarland, J.F.W. McOmie *J. Chem. Soc.* **1951**, 1218-1221; b) F. H. S. Curd, F. C. Rose *J. Chem. Soc.* **1946**, 343-348., 4.2 A (a) Banks *J. Am. Chem. Soc.* **1944**, 66, 1131 b) Ghosh and Dolly *J. Indian Chem. Soc.* **1981**, 58, 512-513.

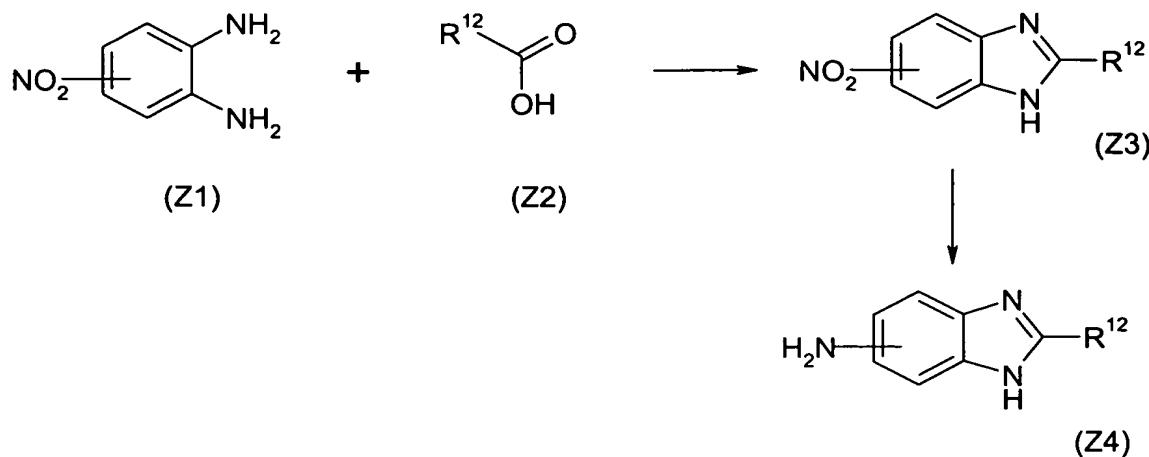
Diagram 4A

For example, in variant 4.1 A, 1 equivalent of the compound (A5) and 1 to 3
 5 equivalents, preferably about 2 equivalents of the compound (A6) are heated
 without a solvent or in an organic solvent such as for example sulpholane,
 dimethylformamide, dimethylacetamide, toluene, N-methylpyrrolidone,
 dimethylsulphoxide or dioxane, preferably sulpholane, for 0.1 to 4 hours,
 preferably 1 hour, at 100 to 220 °C, preferably at about 160 °C. After cooling, the
 10 product (A7) is crystallised by the addition of organic solvents or mixtures of
 solvents, e.g. diethylether/methanol, ethyl acetate, methylene chloride, or
 diethylether, preferably diethylether/methanol 9/1, or purified by chromatography.

For example, in variant 4.2 A , 1 equivalent of the compound (A5) and 1 to 3
 15 equivalents of the compound (A6) are stirred with acid, for example 1-10
 equivalents of 10-38% hydrochloric acid and/or an alcohol, for example ethanol,
 propanol, butanol, preferably ethanol, at reflux temperature for 1 to 48 hours,
 preferably about 5 hours.

The product precipitated (A7) is filtered off and optionally washed with water, dried
 20 and crystallised from a suitable organic solvent.

If R⁶ denotes an optionally substituted benzimidazole, the preparation of the
 compounds (A6) using methods known from the literature may be carried out as
 shown in the following diagram, for example:



Accordingly, for example, 33 mmol of the compound (Z1), 49 mmol of the compound (Z2) and 49 mmol of 1-ethoxycarbonyl-2-ethoxydihydroquinoline

5 (EEDQ) are stirred into 50 ml of an organic solvent, preferably dimethylformamide, at about 100 to 130 °C, preferably at about 115 °C, 1 to 4 hours, preferably about 3 hours. Then the cooled reaction solution is added to 50 to 400 ml, preferably about 200 ml of a water/ethyl acetate mixture (mixing ratio about 1:1). The crystals formed (Z3) are suction filtered and washed.

10 Then 4.2 mmol of the compound (Z3) are stirred with 12.5 mmol of tin(II)chloride and 30 mmol of potassium carbonate in about 50 ml of an organic diluent, preferably ethyl acetate, at about 22 °C for 4 to 48 hours, preferably about 24 hours. After the addition of 22 g of kieselgur the mixture is extracted with an organic diluent or mixture of diluents, preferably with a mixture of dichloromethane

15 / methanol (9:1), the combined extracts are evaporated down and the precipitate formed (Z4) or the crystals produced (Z4) is or are isolated.

Step 5A

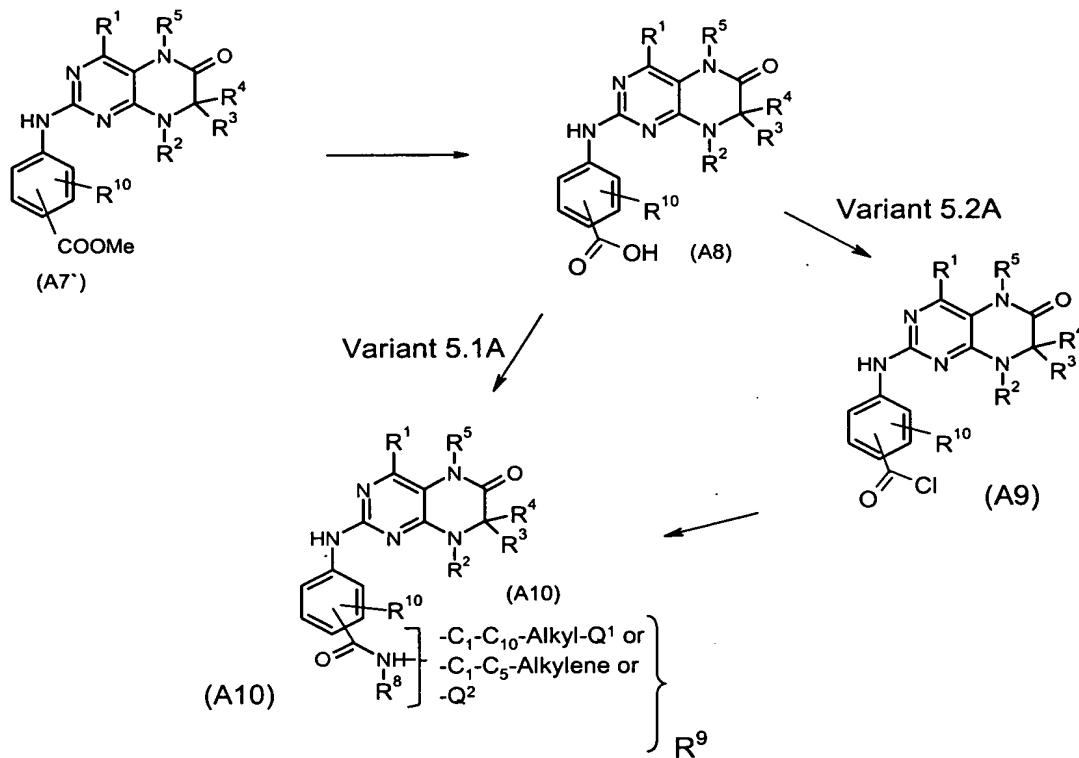
20

If R⁹ denotes -CONR⁸-C₁-C₁₀-alkyl-Q¹, -CONH-C₁-C₅-alkylene or -CONR⁸-Q², wherein the substituents have the meanings given hereinbefore, the compounds according to the invention may be prepared using methods known from the literature, for example as shown in Diagram 5A.

The compound (A7') obtained in Step 4A may be reacted either by saponification and subsequent amination to obtain the amide of general formula (A10) (Diagram (5A) variant 5.1A), or by saponification, with subsequent conversion into the acid chloride (A9) and subsequent amination (Diagram (5A) variant 5.2A).

5

Diagram 5A



10

Variant 5.1 A:

In variant 5.1 A, for example, 20 mmol of the ester (A7') are heated in about 100 ml of a base, preferably 1N sodium hydroxide solution or lithium hydroxide solution and about 500 ml of an alcohol, for example with ethanol, dioxane or methanol, preferably methanol, until the ester is completely reacted. Then the alcohol is distilled off. The residue is taken up in about 200 ml of water and acidified while cooling with acid, for example hydrochloric acid, preferably with 2 N hydrochloric acid. The product (A8) is filtered off and dried.

For example, about 0.5 mmol of the compound (A8) are dissolved with about 0.5 mmol of O-benzotriazolyl-N,N,N',N'-tetramethyluronium tetrafluoroborate (TBTU) and about 1.4 mmol of diisopropylethylamine (DIPEA) in about 5 ml of an organic diluent, for example tetrahydrofuran, dimethylformamide, N-methylpyrrolidone,

5 dimethylacetamide, preferably dimethylformamide. After the addition of about 0.75 mmol of an amine which forms the substituent R⁹, the reaction mixture is stirred for 0.1 to 24 hours, preferably about 12 hours at 20°C to 100°C. The product of formula (A10) is obtained for example by crystallisation or chromatographic purification.

10

Variant 5.2 A:

In variant 5.2 A about 1 mmol of the acid (A8) for example is suspended in about 2.7 ml of thionyl chloride. The mixture is heated to 40°C to 80 °C, preferably about 50 °C, and at constant temperature 2 to 10 drops, preferably about 3 drops of

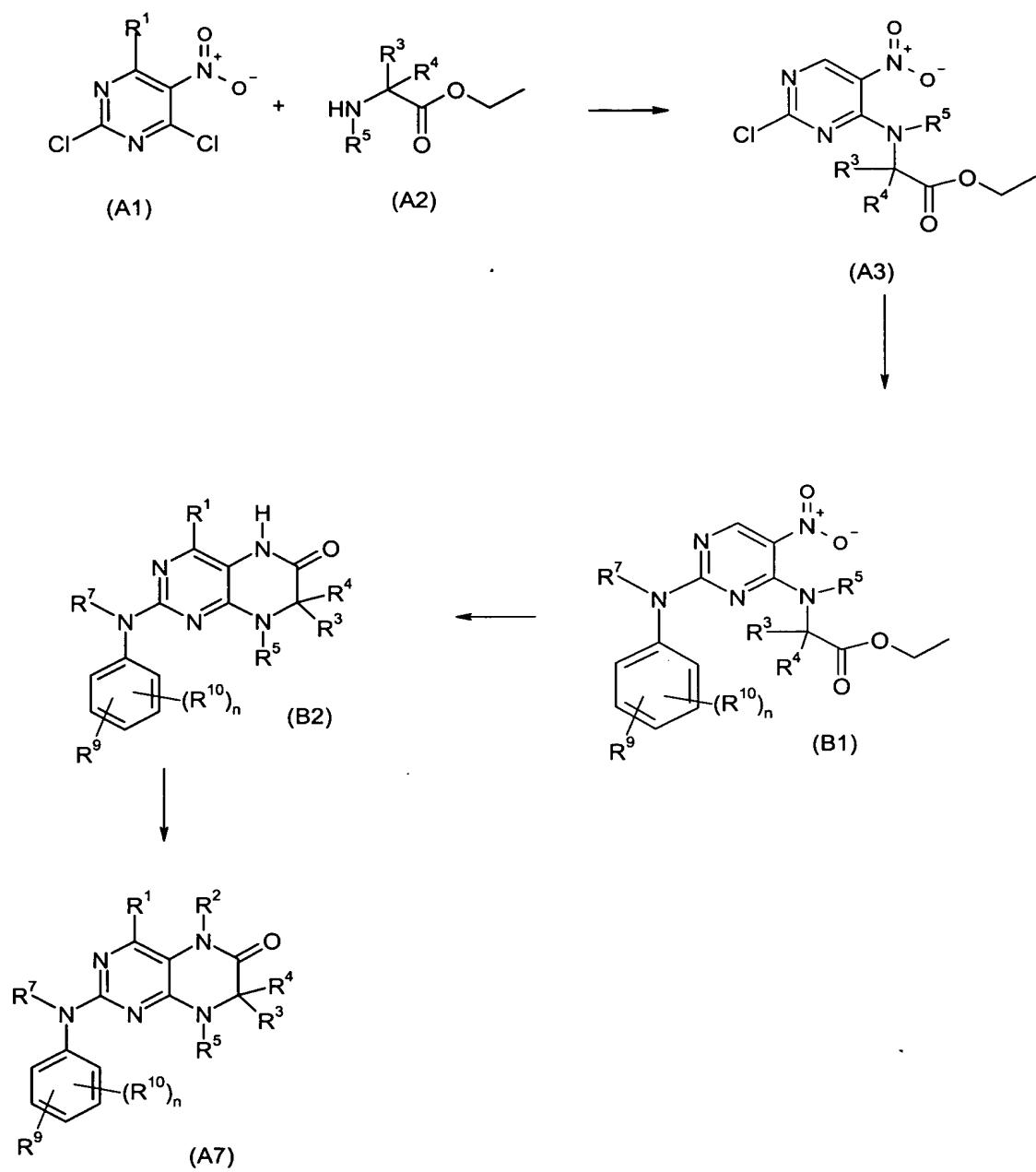
15 dimethylformamide are added to the reaction mixture with stirring. Then stirring is continued at 90°C until the reaction is complete. Excess thionyl chloride is distilled off. About 1 mmol of the acid chloride formed (A9) are dissolved in about 30 ml of an organic diluent, for example dichloromethane. After the addition of an amine which forms the substituent R⁹ the mixture is stirred at about 22°C. The precipitate

20 formed is filtered off and washed with water. The residue remaining is washed with an organic diluent, for example methanol. The mother liquor is purified, for example by chromatography, and evaporated down. The product (A10) remains.

25 Method B

Alternatively to the methods described above, after Step 1A first the compound (A3) may be aminated and then the product (B1) may be cyclised to yield the compound (B2), as shown in Diagram B. Further substitution of the compound

30 (B2) to yield the compound (A7) may be carried out for example as in Step 3A.

Diagram B

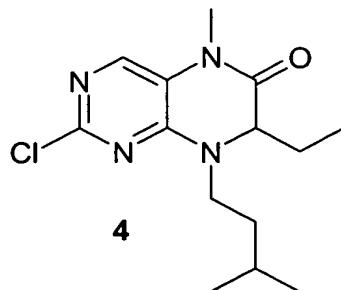
The new compounds of general formula (I) may be synthesised analogously to the following examples of synthesis. These Examples are, however, intended only as examples of procedures to illustrate the invention further, without restricting the invention to their subject matter.

5

Example 63 and Example 109 :

In order to synthesise the compounds 63 and 109, first an intermediate compound 4

10



is prepared as described hereinafter.

38.9 ml (0.263 mol) of ethyl 2-bromobutyrate and 36.4 g (0.263 mol) of potassium 15 carbonate were placed in 350 ml of ethyl acetate, and then 46.7 ml (0.402 mol) of isoamylamine, dissolved in 70 ml of ethyl acetate, were rapidly added dropwise. The mixture was refluxed for 20 h. The salt formed was filtered off, the filtrate was concentrated by evaporation, combined with 50 ml of toluene and again evaporated to dryness.

20 Yield: 54.3 g of a compound 1 (red oil)

54.3 g of compound 1, dissolved in 400 ml acetone, and 30.7 g (0.222 mol) of potassium carbonate were cooled to 8° C with stirring, combined with a solution of 43.1 g (0.222 mol) of 2,4-dichloro-5-nitropyrimidine in 250 ml acetone and then 25 stirred for 24 h at RT.

The suspension formed was concentrated by evaporation, the residue was extracted with water and ethyl acetate, the organic phase was washed with water and NaCl solution, dried over MgSO₄ and evaporated to dryness.

Yield: 87.3 g of a compound **2** (brown oil)

5

44.1 g of compound **2** were dissolved in 800 ml glacial acetic acid and heated to 65°C and 36 g of iron filings were added batchwise. Then the mixture was stirred for 3 h at 70° C, the precipitate was filtered off and the filtrate was concentrated by evaporation.

10 The residue was applied to silica gel in dichloromethane / methanol 90:10, concentrated by evaporation and purified by column chromatography (eluant: ethyl acetate / cyclohexane 1:1).

The residue was precipitated from ethyl acetate / petroleum ether.

Yield: 16.1 g of a compound **3** (beige powder)

15

16.1 g of compound **3** were dissolved in 75 ml of dimethylacetamide and cooled to 5°C under a nitrogen atmosphere with stirring. Then 2.51 g (0.063 mol) of NaH, 60% dispersion in mineral oil, was added, whereupon the temperature temporarily rose to 16°C. After 30 minutes 3.94 ml (0.063 mol) of methyl iodide, dissolved in

20 75 ml dimethylacetamide, were added, and the mixture was stirred for 24 h at 22°C.

The solvent was concentrated by evaporation, combined with 200 ml of water and the precipitate formed was suction filtered, then extracted by stirring with petroleum ether.

25 Yield: 15.1 g of a compound **4** (yellow powder)

¹H-NMR (250 MHz): = 7.80 (1H, s), 4.35 (m, 1H), 3.92 (m, 1H), 3.22 (s, 3H), 3.14 (m, 1H), 1.81 (m, 2H), 1.60- 1.40 (m, 3H), 0.90 (m, 6H), 0.70 (t, 3H).

Synthesis of Example 63

30

2.5 g of compound **4**, 1.43 g of 4-amino-3-methoxybenzoic acid, 1.25 mL of conc. hydrochloric acid, 150 mL of dist. water and 37.5 mL of ethanol were refluxed for 10 h. The precipitate was filtered off, washed with water and extracted by stirring

in methanol. Then the precipitate was recrystallised using petroleum ether and ether.

Yield: 1.6 g of a compound **5** (white powder)

5 0.2 g of compound **5**, 5 mL of benzylamine, 0.16 g of TBTU, 0.17 g of DIPEA were dissolved in 2 ml of dimethylformamide (DMF) and stirred for 48 h at ambient temperature. Then the reaction mixture was taken up in methylene chloride, washed with water and the organic phase was evaporated down. When petroleum ether/ethyl acetate 9:1 was added the product was precipitated in the form of light
10 beige crystals.

Yield: 0.18 g. Melting point: 178°C

Synthesis of Example 109 :

15 5 g of 2 amino-5-nitroaniline, 6.03 g of 4-pyridylcarboxylic acid, 12.1 g of EEDQ are dissolved in 50 mL of DMF and stirred at 115°C for 1.75 h, then the DMF is distilled off *in vacuo* and the reaction mixture is then heated to 180°C for 1 h. The residue is taken up in 30 mL of DMF and combined with 200 mL of water and 100 mL of ethyl acetate. The crystal slurry obtained is filtered off and washed with
20 water, ethyl acetate and ether.

Yield: 5.8 g of a compound **6**

25 2 g of the compound **6** is combined with 0.2 g of 5 % Pd/C in 30 mL of ethanol and hydrogenated in the presence of hydrogen. It is then evaporated down and crystallised from ethanol and toluene.

Yield: 1.75 g of white powder of a compound **7**.

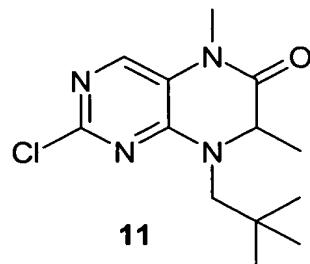
30 0.2 g of the compound **5**, 0.28 g of the compound **7**, 0.001 g of sodium-tert. butoxide, 2.5 mL of ethyleneglycol dimethylether, 0.006 g of palladium(II) acetate and 0.22 g of 2-(di-tert.-butylphospino)biphenyl are dissolved in 1.5 mL of N-methylpyrrolidone (NMP). Then the mixture is heated to 160°C for 0.5 h. The

reaction mixture is then purified over 20 g of silica gel and the product is crystallised from ether, ethyl acetate and petroleum ether.

Yield: 0.04 g of yellow crystals. Melting point: 180°C

5 Example 218, 58 and 4:

In order to synthesise the compounds 218, 58 and 4, first an intermediate compound 11



10 is prepared as described hereinafter.

55.8 g of DL-alanine methyl ester x HCl were dissolved in 500 ml of methanol, then 76.1 ml of 30% sodium methoxide solution was added and the salt was 15 filtered off. 37.8 g of trimethylacetaldehyde were added to the filtrate, then it was left to stand for 22 h. Then 9.5 g of 10% Pd/C was added and the mixture was hydrogenated for 3.1 h at 0.5 bar and 20° C. The reaction mixture was suction filtered through kieselgur and concentrated by evaporation. The residue was taken up in diethylether, the salts were filtered through kieselgur and the filtrate was 20 concentrated by evaporation.

Yield: 55.8 g of a compound 8 (clear liquid)

48.5 g of 2,4-dichloro-5-nitropyrimidine were placed in 400 ml of diethylether, 41.0 g of potassium hydrogen carbonate in 400 ml of water were added, and the 25 mixture was cooled to -5°C. 43.3 g of compound 8 were dissolved in 400 ml of diethylether and added dropwise at -5° C. The mixture was stirred for 1 h at -5°C and for 2 h at 0° C, then heated to ambient temperature and the reaction mixture was left to stand for 24 h.

The organic phase was separated off, dried over MgSO₄ and evaporated to dryness.

Yield: 79.2 g of a compound **9** (yellow resin)

5 79.0 g of compound **9** were dissolved in 1000 ml of glacial acetic acid and heated to 70°C. After the removal of the heat source 52 g of iron was added batchwise. The temperature rose to about 110° C and the mixture was stirred for 1 h at this temperature. The suspension was filtered while hot and the filtrate was concentrated by evaporation.

10 The residue was taken up in ethyl acetate and combined with 150 ml of conc. HCl, the organic phase was separated off and the aqueous phase extracted several times with dichloromethane. The combined organic phases were concentrated by evaporation, applied to silica gel and purified by column chromatography (eluant: petroleum ether/ethyl acetate 1:1).

15 As the isolated substance was still highly contaminated, it was again purified over silica gel. The desired compound crystallised out, the crystals were suction filtered. The mother liquor was concentrated by evaporation and recrystallised from ethyl acetate / diethylether.

Yield: 17.63 g of a compound **10**

20 7.6 g of the compound **10** and 6.4 ml of methyl iodide were placed in 75 ml of dimethylacetamide (DMA) and cooled to –15°C. 1.25 g of NaH, 60% dispersion in mineral oil, was added batchwise, and stirred for 30 min. at –10° to –5° C. Then 150 ml of ice water were added, the crystals were suction filtered and washed with water and petroleum ether. The crystals were taken up in dichloromethane, filtered through kieselgur and the filtrate was evaporated to dryness. It was recrystallised from petroleum ether.

25

Yield: 6.3 g of compound **11** (beige crystals)

¹H-NMR (250 MHz): = 7.73 (1H, s), 4.35 (d, 1H), 4.25 (m, 1H), 3.35 (s, 3H), 2.55

30 (d, 1H), 1.31 (d, 3H), 0.95 (s, 9H).

Synthesis of Example 218

0.2 g of compound **11**, 3,5-difluoro-4-hydroxyaniline and 0.75 mL of sulpholane were heated to 130°C for 15 min, to 140°C for 15 min and to 170°C for 10 min.

5 Then the mixture was combined with ether, the supernatant solution was decanted off and the residue was crystallised from methanol/ether and recrystallised again from methanol.

Yield: 0.15 g of white crystals. Melting point:>250°C

10 Synthesis of Example 4

6.3 g of compound **11** were dissolved in 25 mL of sulpholane at 100°C, then combined with 4.0 g of ethyl 4-aminobenzoate and heated to 170°C for 1 h. Then the mixture was combined with 50 mL of ether. After crystallisation started, a further 50 mL of ether and 50 mL of methanol were added. The crystals were

15 recrystallised from methanol.

Yield: 6.6 g of a compound **12** (yellowish crystals), melting point: from 65°C decomposition sets in

3.55 g of compound **12** were suspended in 250 mL of methanol and at 60°C

20 combined with 25 mL of 4N sodium hydroxide solution. After 6 h, 15 mL of glacial acetic acid were added, the resulting crystals were filtered off and washed with methanol/ether.

Yield: 1.2 g of a compound **13** (white crystals)

25 1.5 g of compound **13** were dissolved in 7.5 mL of thionyl chloride and heated to 80°C for 1 h. Then the thionyl chloride was eliminated by distillation, the residue was stirred with ether, the crystals were suction filtered and washed with ether.

Yield: 1.7 g of a compound **14** (yellow crystals)

30 0.18 g of 3-aminopyridine were dissolved in 10 mL of tetrahydrofuran (THF) and combined with 0.4 mL of triethylamine. Then 0.22 g of compound **14** were added and the mixture was stirred for 16h at ambient temperature. The mixture was

evaporated to dryness, taken up in ethyl acetate, extracted with water, evaporated down again and the product was crystallised from ethyl acetate.

Yield: 0.07 g (beige crystals), Melting point: 215-216°C

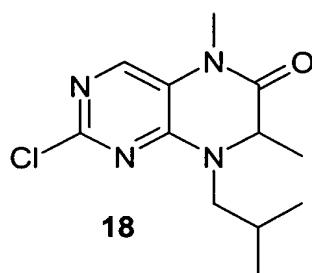
5 Synthesis of Example 58

0.05 g of compound **13** were suspended in 10 mL of dichloromethane, then combined with 0.15 mL of DIPEA and 0.05 g of TBTU. The solution was then stirred for 30 min and combined with 0.01 mL of 4-picollylamine. After 18 h the mixture was combined with 20 mL of water, the organic phase was separated off and the product was purified by silica gel chromatography, then recrystallised from ethyl acetate /petroleum ether.

10 Yield: 0.044 g (white crystals), Melting point: 238-240°C

15 Examples 65 and 125

In order to synthesise the compounds 65 and 125, first an intermediate compound **18**



20 is prepared as described hereinafter.

28.3 g of isobutylamine, 36 g of ethyl R,S-2-bromopropionate and 28 g of potassium carbonate were refluxed in 150 ml of ethyl acetate for 6 h.

After cooling the salt was suction filtered, the mother liquor was concentrated by

25 evaporation.

The residue was combined with 100 ml of toluene and evaporated to dryness.

Yield: 37.2 g of a compound **15** (yellow oil)

38.4 g of 2,4-dichloro-5-nitropyrimidine were placed in 300 ml of diethylether, 30 g of potassium hydrogen carbonate in 300 ml of water were added and the mixture was cooled to 0° C. 37.0 g of compound **15** were dissolved in 300 ml of diethylether and added dropwise at 0°-3° C. After 3 h the phases were separated,

5 the organic phase was dried and evaporated to dryness.

Yield: 71.6 g of a compound **16**

40.0 g of compound **16** were dissolved in 300 ml of glacial acetic acid and heated to 70°C. After removal of the heat source, 30 g of iron was added batchwise. The 10 temperature rose to 110°C. The reaction mixture was cooled to 90°C and stirred for 20 min. at this temperature. Then it was filtered while hot and the filtrate was concentrated by evaporation. The residue was stirred with 300 ml of water and 300 ml of dichloromethane and filtered through kieselgur. The phases were separated. The organic phase was washed with water, dried over MgSO₄ and 15 evaporated to dryness. It was extracted from petroleum ether.

Yield: 26.7 g of a compound **17**

15.0 g of compound **17** were placed in 100 ml of DMA, 4.13 ml of methyl iodide were added and the mixture was cooled to 5° C. 2.60 g of NaH were added 20 batchwise as a 60% dispersion in mineral oil. The temperature rose to 13°C. After 30 min. 300 ml of ice water were added, the crystals precipitated were suction filtered and washed with petroleum ether.

Yield: 13.9 g of a compound **18**

¹H-NMR (250 MHz): = 7.95 (1H, s), 4.30 (m, 1H), 3.95 (m, 1H), 3.24 (s, 3H), 2.95

25 (m, 1H), 2.05 (m, 1H), 1.30 (d, 3H), 0.96 (d, 3H), 0.92 (d, 3H).

Synthesis of Example 65

2.1 g of compound **18** were combined with ethyl 4-aminobenzoate in 10 mL sulpholane and stirred for 2 h at 160°C. Then ether was added and the crystals

30 precipitated were washed with ether:

Yield: 3.0 g of a compound **19**

3 g of the compound **19** were combined with 200 mL of methanol and 25 mL of 4N NaOH and stirred for 4 h at 60°C. Then glacial acetic acid was added, the crystals precipitated were filtered off and washed with ether.

Yield: 2.3 g of a compound **20** (white crystals)

5

0.1 g of compound **20** were suspended in 3 mL of dichloromethane and 3 mL of DMF, and then combined with 0.13 g of DIPEA, 0.095 g of TBTU and 0.045 g of hydroxybenzotriazole (HOBr). Then the solution was stirred for 30 min and combined with 0.035 g of N-methyl-3-picollylamine. After 0.5 h the mixture was combined with water and 1 g of potassium carbonate, the aqueous phase was extracted twice with 50 mL of ethyl acetate and the product was purified by silica gel chromatography and then recrystallised from ethanol/acetone.

10 Yield: 0.08 g

15

Synthesis of Example 125

3.7 g of compound **20**, 3.8 g of TBTU, 1.6 g of HOBr, 5 mL of DIPEA were dissolved in 40 mL of DMF and stirred for 4 h at ambient temperature. The mixture was evaporated down, taken up in 200 mL of ethyl acetate and extracted twice with 5 mL of 5% potassium carbonate solution. The organic phase was evaporated down, the crystals precipitated were filtered off and washed with ethyl acetate and ether.

Yield: 1.65 g of a compound **21** (yellowish crystals)

25 0.486 g of compound **21** were refluxed with 0.33 g of 1,2- phenylenediamine in 10 mL of toluene for 0.5 h, then the mixture was evaporated down. The residue was combined with 100 mL ethyl acetate, the organic phase was extracted twice with water. The organic phase was evaporated down, the crystals precipitated were suction filtered and washed with a little ethyl acetate.

30 Yield: 0.25 g of a compound **22** (white crystals)

0.22 g of compound **22** were stirred into 20 g of polyphosphoric acid for 0.5 h at 150°C, then the mixture was poured onto ice and ammonia was added. It was then extracted twice with 100 mL of ethyl acetate, the organic phase was washed with water and evaporated down. The precipitated product (crystals) was suction

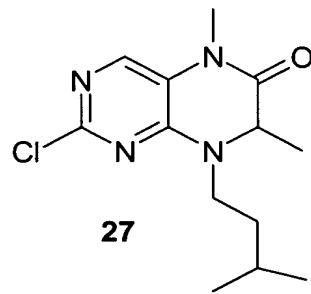
5 filtered and washed with ethyl acetate and ether.

Yield: 0.115 g of yellowish crystals, Melting point: 287°C (decomposition)

Example 171

10

In order to synthesise compound **171** first an intermediate compound **27** [sic]



15

34.4 g of N-isopentyl-benzylamine, 36.2 g of ethyl 2-bromo-propionate and 42.0 g of potassium carbonate were placed in 250 ml of DMF and stirred for 3 h at 110°C. After cooling the inorganic salts were filtered off, the filtrate was concentrated by evaporation. The residue was extracted with water and diethylether, the organic

20 phase was washed with water, dried and evaporated to dryness.

Yield: 55.5 g of a compound **23**

55.5 g of compound **23** were placed in 600 ml of ethanol, and hydrogenated with 25 20 ml of 32% HCl and 6 g of 10% Pd/C at 20°C under 5 bar for 20 min. Then it was filtered through kieselgur and concentrated by evaporation. The residue was combined with 400 ml of diethylether, the precipitate was suction filtered and washed with diethylether.

Yield: 23.5 g of a compound **24**, melting point 105°C

23.5 g of compound **24** were dissolved in 200 ml of water and combined with 20.0 g (0.103 mol) of 2,4 -dichloro-5-nitropyrimidine in 400 ml of diethylether. After the reaction mixture had been cooled to -10° C , 50.0 g (0.499 mol) of potassium carbonate were added batchwise. The mixture was stirred at -5°C for 1 h and at 0°C for 1 h, then heated to ambient temperature. The aqueous phase was separated off, the organic phase was washed with water, dried and evaporated to dryness.

10 Yield: 36.9 g of a compound **25**

20.0 g of the compound **25** were dissolved in 280 ml of glacial acetic acid and heated to 70°C. After removal of the heat source 17 g of iron were added. The temperature rose to 100°C, then the mixture was stirred for 30 min. at this 15 temperature.

It was then filtered while hot and the filtrate was concentrated by evaporation. The residue was combined with 300 ml of dichloromethane and 30 ml of 32% HCl, the phases were separated, the aqueous phase was extracted with dichloromethane, the combined organic phases were washed with water and aqueous ammonia 20 solution, dried and evaporated to dryness. The residue was extracted with diethylether.

Yield: 10.5 g of a compound **26**, melting point: 182°-185° C

2.7 g of the compound **26** and 2.5 ml of methyl iodide were placed in 27 ml of DMA and cooled to -10° C. 0.45 g of NaH, 60% dispersion in mineral oil, was added and stirred for 30 min. at -5°C. Then 10 g of ice and 5 ml of 2N HCl were added and the mixture was concentrated by evaporation. The residue was extracted with ethyl acetate and water, the organic phase was dried, evaporated to dryness and filtered through silica gel.

30 Yield: 3.0 g of compound **27** (oil)

¹H-NMR (250 MHz): = 7.67 (1H, s), 4.32-4.07 (m, 2H), 3.32 (s, 3H), 3.08 (m, 1H), 1.70-1.50 (m, 3H), 1.42 (d, 3H), 0.95 (m, 6H).

Synthesis of Example 171

0.28 g of compound **27**, 0.9 mL of sulpholane and 0.22 g of p-aminobenzoic acid-benzylamide were stirred for 0.5 h at 170°C, then the mixture was combined with

5 ether and the crystals were filtered off. The product was recrystallised from ethanol.

Yield: 0,15 g, melting point: 228-240°C (yellowish crystals)

The compounds of formula (I) listed in Table 1 are obtained analogously to the
10 process described above.

The abbreviations X_2 , X_3 , X_4 , X_5 and X_6 used in Table 1 in each case denote a link to a position in the general formula shown under Table 1 instead of the corresponding groups R^2 , R^3 , R^4 , R^5 and R^6 .

Table 1 (Continued)

Ex.	$\begin{array}{c} R^2 \\ \\ X_2 \\ \backslash \\ CH_3 \end{array}$	R^3	R^4	config. R^3 or R^4	R^5	R^8	mp. [°C]
1	$\begin{array}{c} X_2 \\ \\ CH_3 \end{array}$	H	H	rac.			
2	$\begin{array}{c} CH_3 \\ \\ X_2 \end{array}$	$X_3 \backslash CH_3$	H	rac.			208
3	$\begin{array}{c} CH_3 \\ \\ X_2 \end{array}$	$X_3 \backslash CH_3$	H	rac.			241
4	$\begin{array}{c} X_2 \\ \\ CH_3 \end{array}$	$CH_3 \backslash X_3$	H	rac.			
5	$\begin{array}{c} CH_3 \\ \\ X_2 \end{array}$	$X_3 \backslash CH_3$	H	rac.			175
6	$\begin{array}{c} X_2 \\ \\ CH_3 \end{array}$	$CH_3 \backslash X_3$	H	R			190
7	$\begin{array}{c} X_2 \\ \\ CH_3 \end{array}$	$CH_3 \backslash CH_2 X_3$	H	rac.			

Table 1 (Continued)

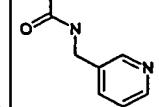
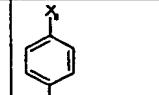
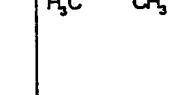
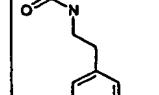
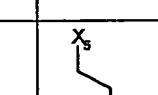
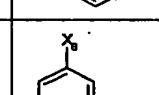
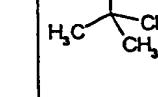
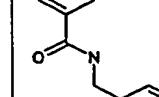
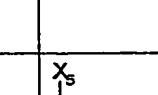
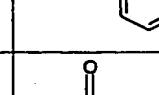
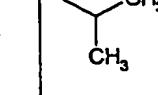
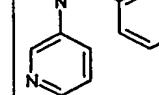
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
8			H	rac.			200
9			H	rac.			168
10			H	rac.			190
11			H	rac.			
12			H	rac.			
13			H	rac.			145
14			H	rac.			

Table 1 (Continued)

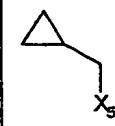
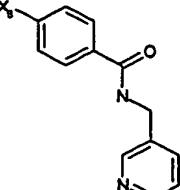
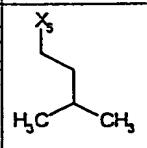
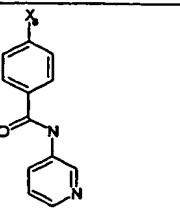
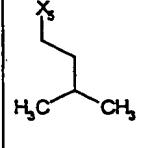
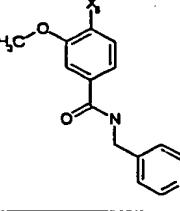
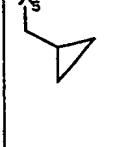
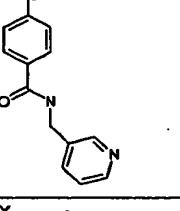
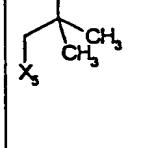
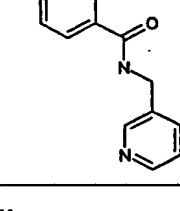
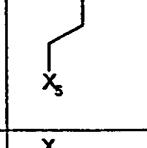
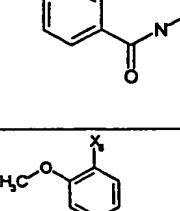
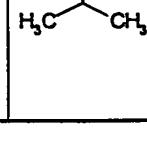
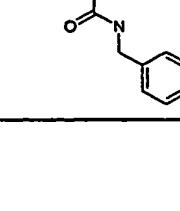
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
15	$\begin{array}{c} X_2 \\ \\ CH_3 \end{array}$	H_3C-X_3	H	rac.			55
16	$\begin{array}{c} CH_3 \\ \\ X_2 \end{array}$	X_3-X_3	H	rac.			250
17	$\begin{array}{c} CH_3 \\ \\ X_2 \end{array}$	X_3-X_3	H	rac.			204
18	$\begin{array}{c} CH_3 \\ \\ X_2 \end{array}$	X_3-X_3	H	rac.			
19	$\begin{array}{c} X_2 \\ \\ CH_3 \end{array}$	H_3C-X_3	H	rac.			
20	$\begin{array}{c} X_2 \\ \\ CH_3 \end{array}$	H_3C-X_3	H	R			221
21	$\begin{array}{c} CH_3 \\ \\ X_2 \end{array}$	X_3-X_3	H	rac.			172

Table 1 (Continued)

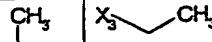
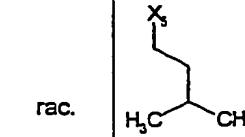
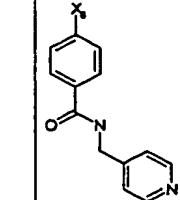
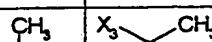
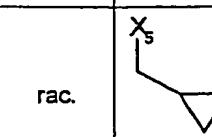
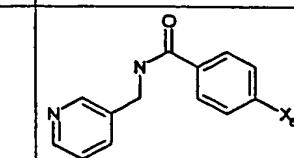
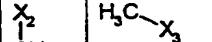
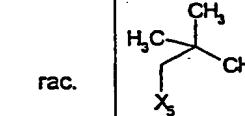
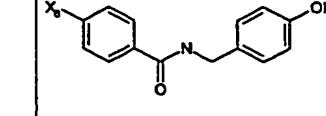
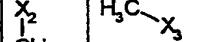
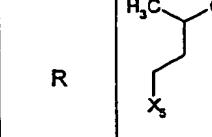
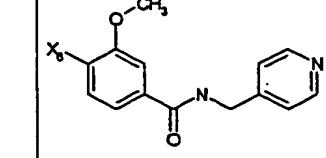
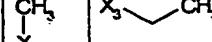
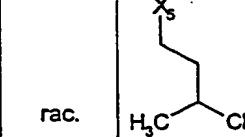
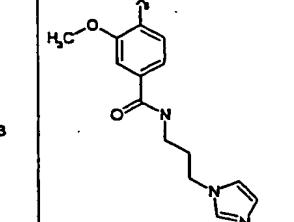
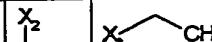
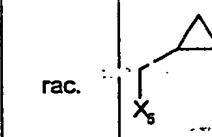
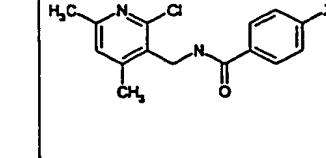
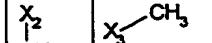
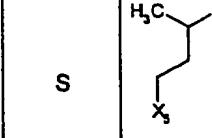
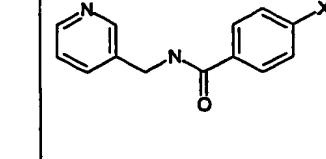
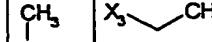
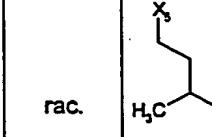
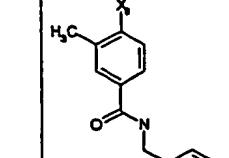
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24				H	rac.			210
25				H	R			213
26				H	rac.			188
27				H	rac.			
28				H	S			
29				H	rac.			178

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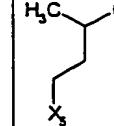
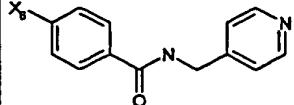
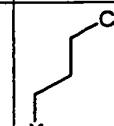
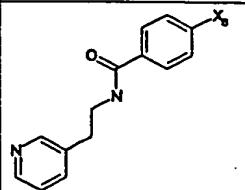
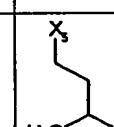
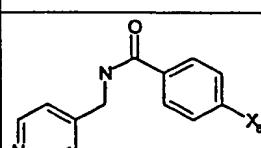
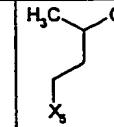
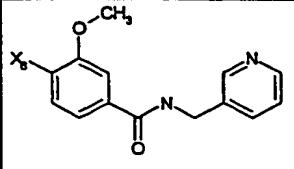
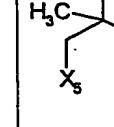
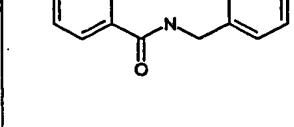
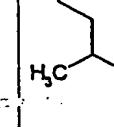
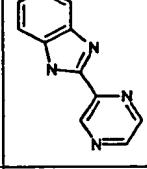
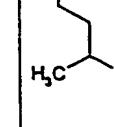
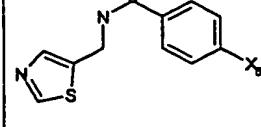
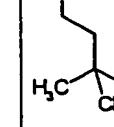
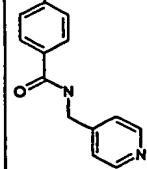
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp.[°C]	
30				H	R			175
31				H	rac.			
32				H	rac.			221
33				H	R			124
34				H	rac.			136
35				H	rac.			162
36				H	rac.			169
37				H	rac.			219

Table 1 (Continued)

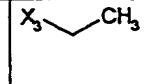
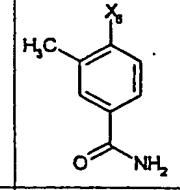
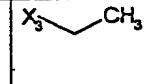
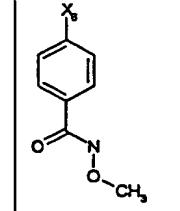
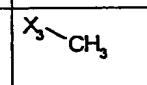
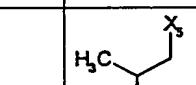
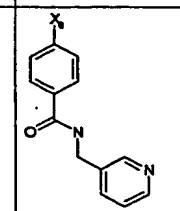
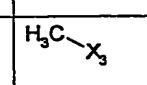
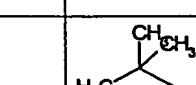
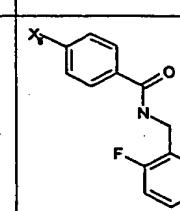
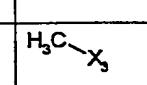
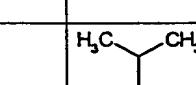
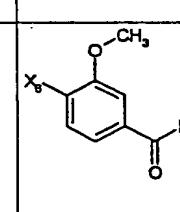
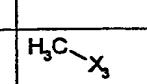
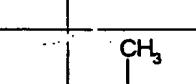
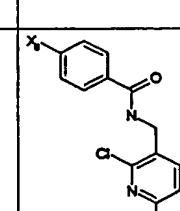
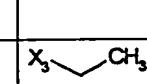
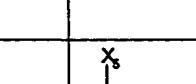
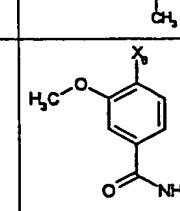
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]	
38				H	rac.			179
39				H	rac.			211
40				H	rac.			
41				H	rac.			
42				H	R			100
43				H	rac.			175
44				H	rac.			203

Table 1 (Continued)

Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
45			H	rac.			165
46			H	rac.			
47			H	rac.			
48			H	rac.			
49			H	rac.			
50			H	rac.			212
51			H	S			
52			H	rac.			

Table 1 (Continued)

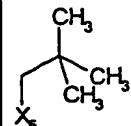
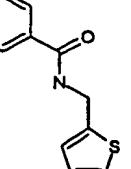
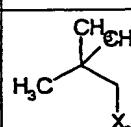
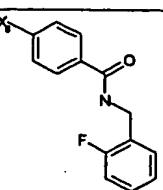
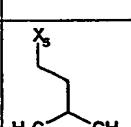
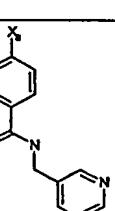
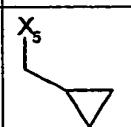
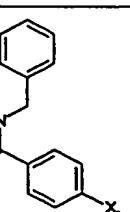
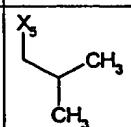
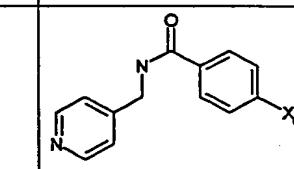
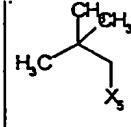
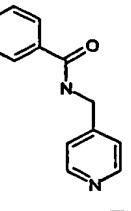
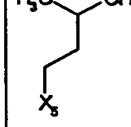
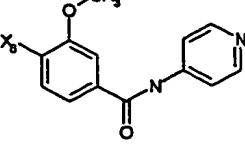
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp.[°C]
53	X ₂ CH ₃	H ₃ C—X ₃	H	rac.			
54	X ₂ CH ₃	H ₃ C—X ₃	H	rac.			
55	CH ₃ X ₂	X ₃ —CH ₃ —CH ₃	H	rac.			191
56	CH ₃ X ₂	X ₃ —CH ₃	H	rac.			158
57	CH ₃ X ₂	X ₃ —CH ₃	H	rac.			230
58	X ₂ CH ₃	H ₃ C—X ₃	H	rac.			
59	X ₂ CH ₃	H ₃ C—X ₃	H	R			125

Table 1 (Continued)

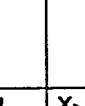
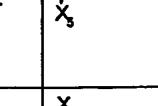
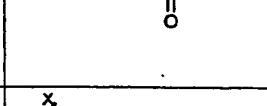
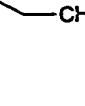
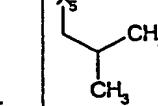
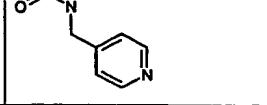
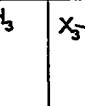
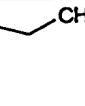
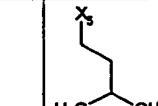
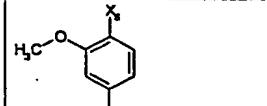
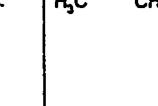
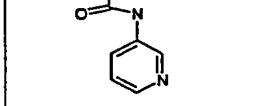
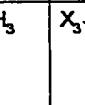
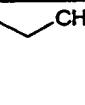
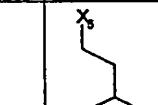
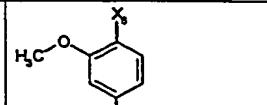
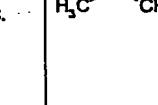
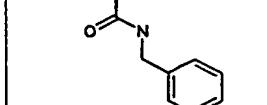
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
60		H	H	rac.			250
61			H	rac.			
62			H	rac.			169
63			H	rac.			178
64			H	rac.			
65			H	rac.			
66			H	R			225

Table 1 (Continued)

Table 1 (Continued)

Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
74	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$				$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C} - \text{CH}_2 - \text{CH}_3 \end{array}$		167
75	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} - \text{X}_3 \\ \quad \\ \text{X}_4 \quad \text{H}_3\text{C} \end{array}$	H	rac.	$\begin{array}{c} \text{CH}_3 - \text{CH}_2 - \text{CH}(\text{CH}_3) - \text{X}_5 \\ \\ \text{H}_3\text{C} \end{array}$		
76	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 - \text{CH}_3 \\ \quad \\ \text{H}_3\text{C} \quad \text{X}_4 \end{array}$	H	rac.	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C} - \text{CH}_2 - \text{CH}(\text{CH}_3) - \text{CH}_3 \end{array}$		246
77	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} - \text{X}_3 \\ \quad \\ \text{X}_4 \quad \text{H}_3\text{C} \end{array}$	H	rac.	$\begin{array}{c} \text{CH}_3 - \text{CH}(\text{CH}_3) - \text{CH}(\text{CH}_3) - \text{X}_5 \\ \\ \text{H}_3\text{C} \end{array}$		
78	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 - \text{CH}_3 \\ \quad \\ \text{H}_3\text{C} \quad \text{X}_4 \end{array}$	H	rac.	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C} - \text{CH}_2 - \text{CH}(\text{CH}_3) - \text{CH}_3 \end{array}$		172
79	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 - \text{CH}_3 \\ \quad \\ \text{H}_3\text{C} \quad \text{CH}_3 \end{array}$	H	rac.	$\begin{array}{c} \text{H}_3\text{C} - \text{CH}(\text{CH}_3) - \text{CH}(\text{CH}_3) - \text{X}_5 \\ \\ \text{H}_3\text{C} \end{array}$		170
80	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 - \text{CH}_3 \\ \quad \\ \text{H}_3\text{C} \quad \text{CH}_3 \end{array}$	$\begin{array}{c} \text{X}_4 - \text{CH}_3 \\ \quad \\ \text{H}_3\text{C} \quad \text{H}_3\text{C} \end{array}$	rac.	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C} - \text{CH}_2 - \text{CH}_3 \end{array}$		222
81	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 - \text{CH}_3 \\ \quad \\ \text{H}_3\text{C} \quad \text{CH}_3 \end{array}$	H	rac.	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C} - \text{CH}_2 - \text{CH}_3 \end{array}$		187

Table 1 (Continued)

Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
82			H	rac.			215
83							199
84				rac.			127
85			H	rac.			
86			H	rac.			169
87			H	rac.			250
88			H	rac.			233
89			H	rac.			160

Table 1 (Continued)

Ex.	$R^2 - -$	R^3	R^4	config. R^3 or R^4	R^5	R^6	mp. [°C]
90	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$	H	rac.	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \\ \\ \text{CH}_3 \end{array}$		154
91	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{CH}_2\text{CH}_2 \\ \\ \text{X}_3 \end{array}$	H	rac.			
92	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_2\text{CH}_3 \end{array}$	H	rac.	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_2\text{CH}_2 \\ \\ \text{H}_3\text{C} \\ \\ \text{CH}_3 \end{array}$		
93	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	rac.	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{CH}_2\text{CH}_2 \\ \\ \text{CH}_3 \\ \\ \text{CH}_3 \end{array}$		
94	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	R	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{CH}_2\text{CH}_2 \\ \\ \text{X}_3 \end{array}$		
95	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_2\text{CH}_3 \end{array}$	H	rac.	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_2\text{CH}_2 \\ \\ \text{CH}_3 \end{array}$		150
96	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{X}_4 \\ \\ \text{CH}_3 \end{array}$	rac.	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_2\text{CH}_2 \\ \\ \text{H}_3\text{C} \\ \\ \text{CH}_3 \end{array}$		300
97	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	rac.	$\begin{array}{c} \text{CH}_3 \\ \\ \text{CH}_2\text{CH}_2 \\ \\ \text{CH}_3 \\ \\ \text{CH}_3 \end{array}$		243

Table 1 (Continued)

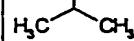
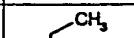
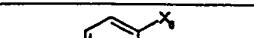
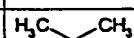
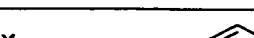
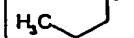
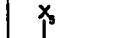
Ex	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
98	CH ₃ X ₂	X ₃ —CH ₃	H	rac.			209
99	CH ₃ X ₂	X ₃ —CH ₃	H	rac.			182
100	X ₂ CH ₃	X ₃ —CH ₃	H	rac.			
101	X ₂ CH ₃	X ₃ —CH ₃	H	R			232
102	CH ₃ X ₂	X ₃ —CH ₃	H	rac.			
103	CH ₃ X ₂	X ₃ —CH ₃	H	rac.			
104	CH ₃ X ₂	X ₃ —CH ₃	H	rac.			146

Table 1 (Continued)

Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
105				H rac.			209
106				H rac.			286
107				H rac.			
108				H R			202
109				H rac.			180
110				H rac.			
111				H rac.			250

Table 1 (Continued)

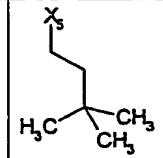
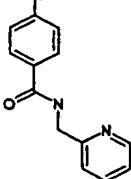
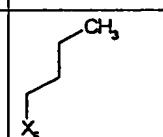
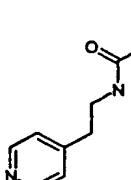
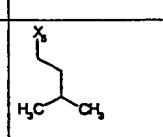
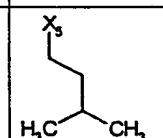
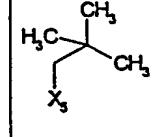
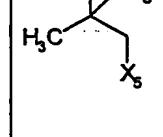
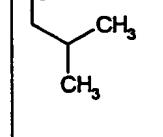
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
112	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$		H	rac.		
113	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$		H	rac.		
114	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{X}_4 \\ \\ \text{CH}_3 \end{array}$				237
115	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$		H	rac.		135
116	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$		H	rac.		
117	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$		H	rac.		
118	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$		H	rac.		

Table 1 (Continued)

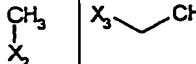
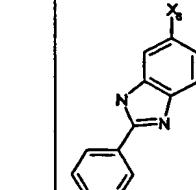
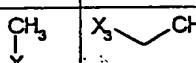
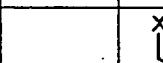
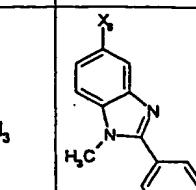
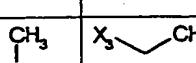
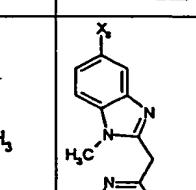
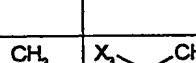
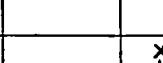
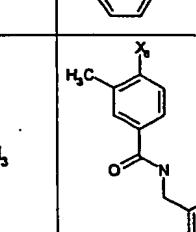
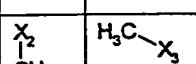
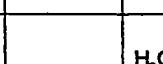
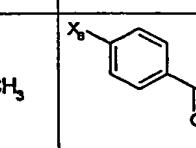
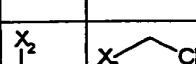
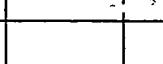
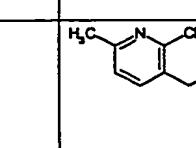
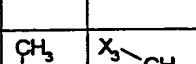
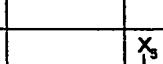
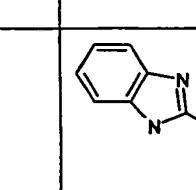
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]	
119				H	rac.			213
120				H	rac.			198
121				H	rac.			
122				H	rac.			
123				H	rac.			
124				H	rac.			
125				H	rac.			287

Table 1 (Continued)

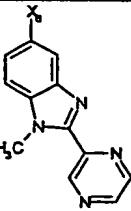
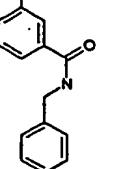
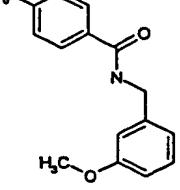
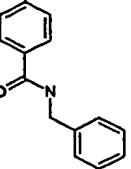
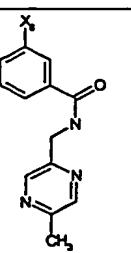
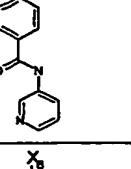
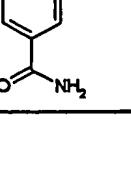
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp.[°C]
126	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$		H rac.	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C} - \text{CH}_2 - \text{CH}_3 \end{array}$		195
127	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$		H rac.	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C} - \text{CH}_2 - \text{CH}_3 \end{array}$		
128	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$		H rac.	$\begin{array}{c} \text{CH}_2\text{CH}_3 \\ \\ \text{H}_3\text{C} - \text{CH}_2 - \text{CH}_2 - \text{X}_5 \end{array}$		
129	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$		H rac.	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C} - \text{CH}_2 - \text{CH}(\text{CH}_3)_2 \end{array}$		247
130	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$		H rac.	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C} - \text{CH}_2 - \text{CH}_3 \end{array}$		
131	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{X}_4 \\ \\ \text{CH}_3 \end{array}$		$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C} - \text{CH}_2 - \text{CH}_3 \end{array}$		281
132	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$		H rac.	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{H}_3\text{C} - \text{CH}_2 - \text{CH}_2 - \text{X}_5 \end{array}$		

Table 1 (Continued)

Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp.[°C]
133	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \text{---} \text{CH}_3 \\ \\ \text{X}_3 \end{array}$	H	rac.			
134	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \text{---} \text{CH}_3 \\ \\ \text{X}_3 \end{array}$	H	rac.			208
135	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{X}_3 \text{---} \text{CH}_3 \\ \\ \text{X}_3 \end{array}$	H	rac.			
136	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{X}_3 \text{---} \text{CH}_3 \\ \\ \text{X}_3 \end{array}$	H	R			192
137	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{X}_3 \text{---} \text{CH}_3 \\ \\ \text{X}_3 \end{array}$	H	rac.			212
138	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \text{---} \text{X}_3 \\ \\ \text{X}_3 \end{array}$	H	rac.			
139	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \text{---} \text{CH}_3 \\ \\ \text{X}_3 \end{array}$	H	rac.			

Table 1 (Continued)

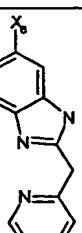
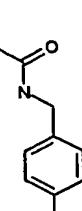
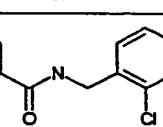
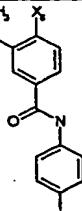
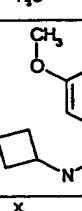
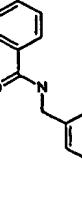
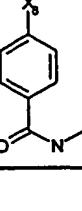
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
140	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	X_3-CH_3	H	rac.	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \end{array}$		148
141	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	rac.	$\begin{array}{c} \text{CH}_3-\text{CH}_3 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \\ \\ \text{X}_5 \end{array}$		
142	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	rac.	$\begin{array}{c} \text{CH}_3 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \\ \\ \text{X}_5 \end{array}$		
143	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	X_3-CH_3	H	rac.	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \\ \\ \text{CH}_3 \end{array}$		186
144	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{X}_4 \\ \\ \text{C}_3 \end{array}$			$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \end{array}$		199
145	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	X_3-CH_3	H	rac.	$\begin{array}{c} \text{X}_5 \\ \\ \text{C}_3 \end{array}$		214
146	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	X_3-CH_3	H	rac.	$\begin{array}{c} \text{X}_5 \\ \\ \text{C}_3 \end{array}$		155

Table 1 (Continued)

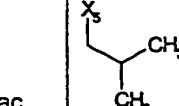
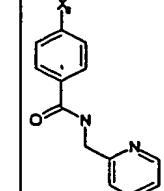
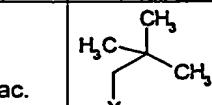
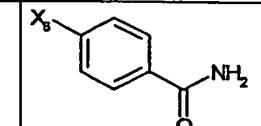
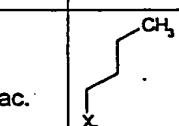
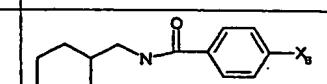
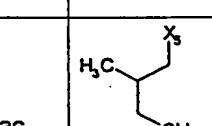
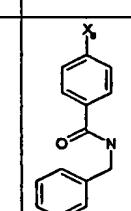
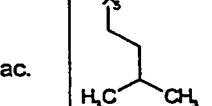
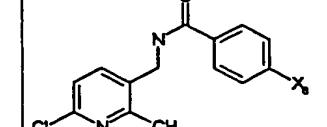
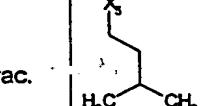
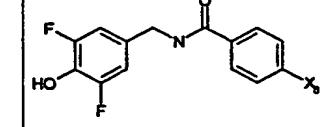
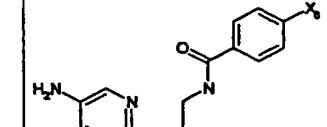
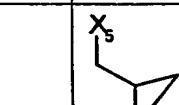
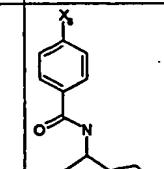
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
147	CH ₃ X ₂	X ₃ —CH ₃	H	rac.			
148	X ₂ CH ₃	H ₃ C—X ₃	H	rac.			
149	X ₂ CH ₃	X ₃ —CH ₃	H	rac.			245
150	CH ₃ X ₂	X ₃ —CH ₃	H	rac.			
151	CH ₃ X ₂	X ₃ —CH ₃	H	rac.			
152	CH ₃ X ₂	X ₃ —CH ₃	H	rac.			
153	X ₂ CH ₃	X ₃ —CH ₃	H	rac.			
154	CH ₃ X ₂	X ₃ —CH ₃	H	rac.			

Table 1 (Continued)

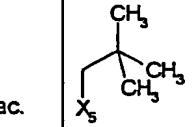
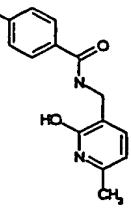
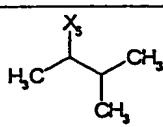
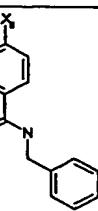
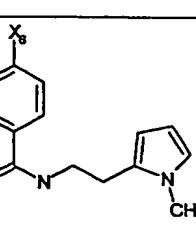
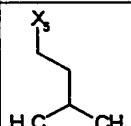
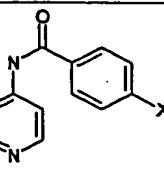
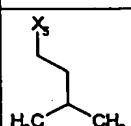
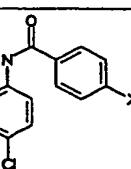
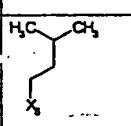
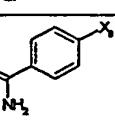
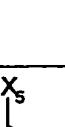
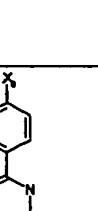
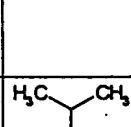
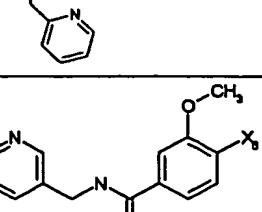
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp.[°C]	
155				H				
156				H			265	
157				H			192	
158				H			222	
159				H			221	
160							298	
161				H			181	
162				H	S			

Table 1 (Continued)

Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
163	CH ₃ X ₂		X ₃ — X ₄		X ₅ H ₃ C—CH ₃	CH ₃ O— X ₆ C ₆ H ₄ — C(=O)N— C ₆ H ₄ — CH ₃	172
164	CH ₃ X ₂	X ₃ —CH ₃	H	rac.	X ₅ H ₃ C— CH ₃	CH ₃ O— C ₆ H ₅ C(=O)N— CH ₃ — CH ₃	227
165	CH ₃ X ₂	X ₃ —CH ₃	H	rac.	X ₅ H ₃ C—CH ₃	CH ₃ C ₆ H ₅ — C ₆ H ₄ — OH— N— C ₆ H ₄ — C(=O)O— C ₆ H ₅	258
166	CH ₃ X ₂	X ₃ —CH ₃	X ₄ — CH ₃		X ₅ H ₃ C—CH ₃	CH ₃ O— C ₆ H ₅ C(=O)N— CH ₃	266
167	X ₂ CH ₃	H ₃ C—X ₃	H	rac.	CH ₃ — CH ₂ — CH ₃ X ₅	CH ₃ O— C ₆ H ₅ C(=O)N— CH ₂ — C ₆ H ₅ C(=O)NH ₂	
168	CH ₃ X ₂	X ₃ —CH ₃	X ₄ — CH ₃	rac.	X ₅ H ₃ C—CH ₃	CH ₃ O— C ₆ H ₅ C(=O)O— C ₆ H ₄ — C(=O)O— C ₆ H ₅	159
169	CH ₃ X ₂		X ₃ — X ₄		X ₅ H ₃ C—CH ₃	CH ₃ O— X ₆ C ₆ H ₄ — C(=O)N— C ₆ H ₄ — CH ₃	250

Table 1 (Continued)

Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
170			H	rac.			213
171			H	rac.			228
172			H	rac.			181
173			H	rac.			182
174			H	rac.			
175			H	rac.			197
176			H	rac.			

Table 1 (Continued)

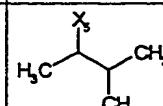
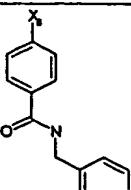
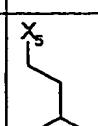
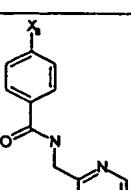
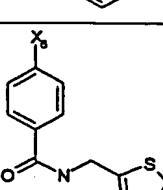
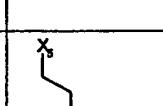
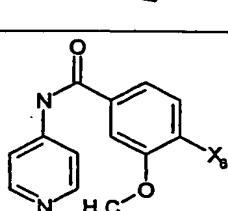
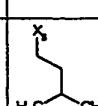
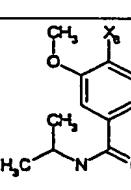
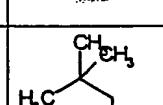
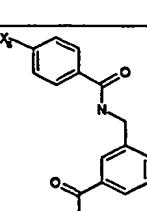
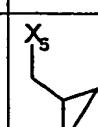
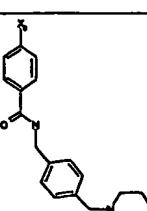
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp.[°C]
177	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{X}_3 \text{---} \text{CH}_3$	H	rac.			216
178	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{X}_3 \text{---} \text{CH}_3$	H	rac.			200
179	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{X}_3 \text{---} \text{CH}_3$	H	rac.			197
180	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{X}_3 \text{---} \text{CH}_3$	$\text{X}_4 \text{---} \text{CH}_3$	rac.			143
181	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$						234
182	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C} \text{---} \text{X}_3$	H	rac.			
183	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{X}_3 \text{---} \text{CH}_3$	H	rac.			169

Table 1 (Continued)

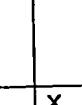
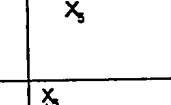
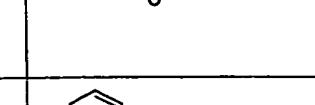
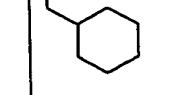
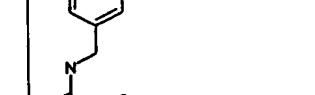
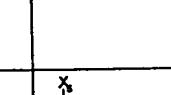
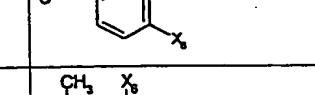
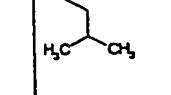
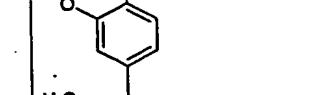
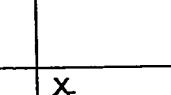
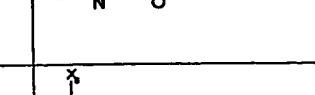
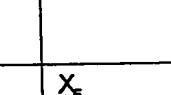
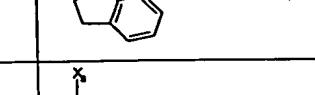
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
184				H	rac.		
185				H	rac.		
186							
187				H	rac.		
188				H	rac.		
189				H	rac.		
190							

Table 1 (Continued)

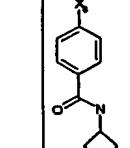
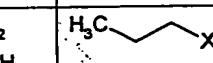
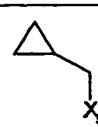
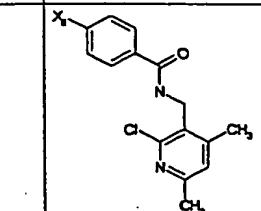
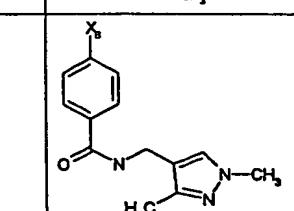
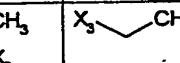
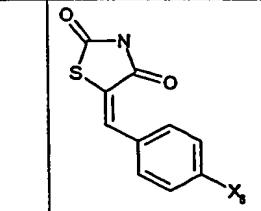
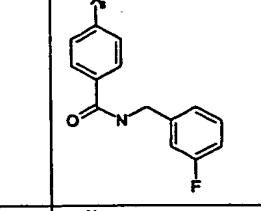
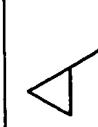
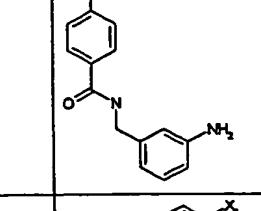
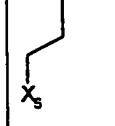
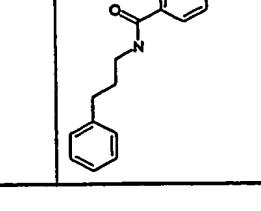
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
191							253
192			H	rac.			
193			H	rac.			201
194			H	rac.			250
195			H	rac.			198
196			H	rac.			245
197			H	rac.			

Table 1 (Continued)

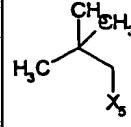
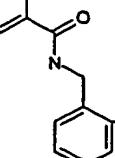
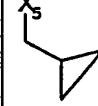
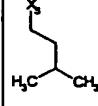
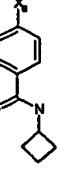
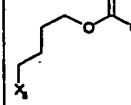
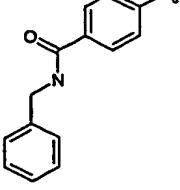
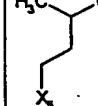
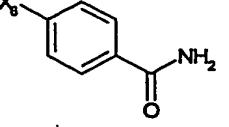
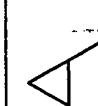
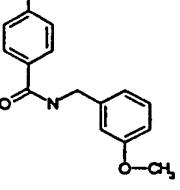
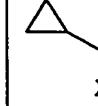
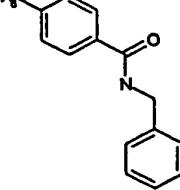
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
198	$\begin{matrix} X_2 \\ \\ CH_3 \end{matrix}$	H_3C-X_3	H	rac.			
199	$\begin{matrix} CH_3 \\ \\ X_2 \end{matrix}$	X_3-X_3	H	rac.			
200	$\begin{matrix} CH_3 \\ \\ X_2 \end{matrix}$	X_3-CH_3	$\begin{matrix} X_4 \\ \\ CH_3 \end{matrix}$				198
201	$\begin{matrix} X_2 \\ \\ CH_3 \end{matrix}$	H	H	rac.			
202	$\begin{matrix} X_2 \\ \\ CH_3 \end{matrix}$	H_3C-X_3	H	rac.			
203	$\begin{matrix} CH_3 \\ \\ X_2 \end{matrix}$	X_3-CH_3	H	rac.			198
204	$\begin{matrix} X_2 \\ \\ CH_3 \end{matrix}$	H_3C-X_3	H	rac.			

Table 1 (Continued)

Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
205	X ₂ CH ₃	H ₃ C— X ₃	H	rac.	△— X ₅	X ₆ — C ₆ H ₄ — C(=O)N— C ₆ H ₄ — Cl	
206	X ₂ CH ₃	H ₃ C— X ₃	H	rac.	△— X ₅	X ₆ — C ₆ H ₄ — O—CH ₃ C(=O)N— C ₆ H ₄ — C ₆ H ₅	
207	CH ₃ X ₂	X ₃ —CH ₃	H	rac.	△— X ₅	X ₆ — C ₆ H ₄ — C(=O)N— C ₆ H ₄ — NH ₂	184
208	X ₂ CH ₃	X ₃ —CH ₃	H	rac.	CH ₃ — X ₅	— C ₆ H ₅ — C ₆ H ₄ — N— C ₆ H ₅ — X ₆	253
209	CH ₃ X ₂	X ₃ —CH ₃	H	rac.	X ₅ — △	O—CH ₃ C(=O)C ₆ H ₄ — X ₆	240
210	X ₂ CH ₃	H ₃ C— X ₃	H	rac.	CH ₃ — CH ₃ H ₃ C— X ₅	X ₆ — C ₆ H ₄ — C(=O)N— C ₆ H ₄ — C(F)(F) ₂	
211	CH ₃ X ₂	X ₃ —CH ₃	X ₄ CH ₃		X ₅ — H ₃ C— CH ₃	X ₆ — C ₆ H ₄ — C(=O)N— CH ₃	266

Table 1 (Continued)

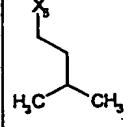
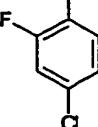
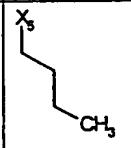
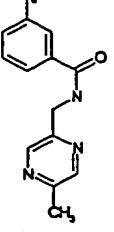
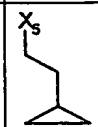
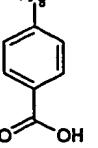
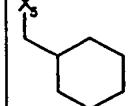
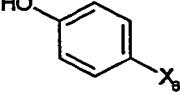
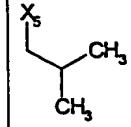
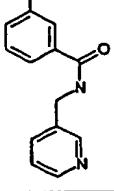
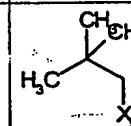
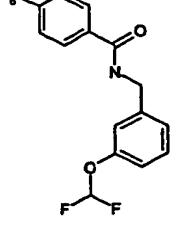
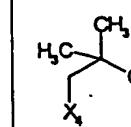
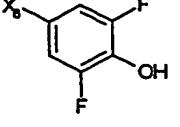
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp.[°C]
212	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$	H	rac.			
213	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$	H	rac.			
214	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$	H	rac.			
215	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$	H	rac.			232
216	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$	H	rac.			
217	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	rac.			
218	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	rac.			>250

Table 1 (Continued)

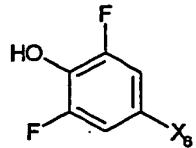
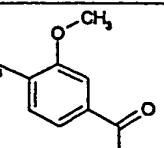
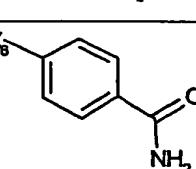
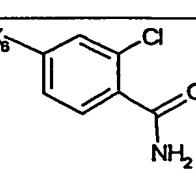
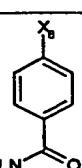
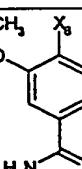
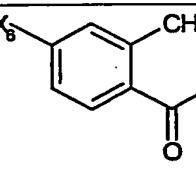
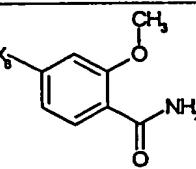
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp.[°C]
219	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	rac.	$\begin{array}{c} \text{X}_3 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \end{array}$		260 (Zers.)
220	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	$\begin{array}{c} \text{H}_3\text{C}-\text{CH}_3 \\ \\ \text{X}_3 \end{array}$		190
221	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	$\begin{array}{c} \text{H}_3\text{C}-\text{CH}_3 \\ \\ \text{X}_3 \end{array}$		228
222	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	$\begin{array}{c} \text{H}_3\text{C}-\text{CH}_3 \\ \\ \text{X}_3 \end{array}$		
223	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	X_3-CH_3	H	R	$\begin{array}{c} \text{X}_3 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \end{array}$		243
224	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	X_3-CH_3	H	R	$\begin{array}{c} \text{X}_3 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \end{array}$		258
225	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	$\begin{array}{c} \text{H}_3\text{C}-\text{CH}_3 \\ \\ \text{X}_3 \end{array}$		
226	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	$\begin{array}{c} \text{H}_3\text{C}-\text{CH}_3 \\ \\ \text{X}_3 \end{array}$		

Table 1 (Continued)

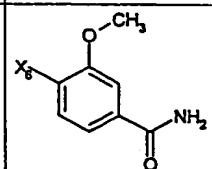
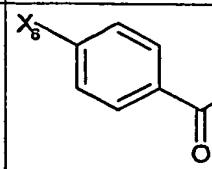
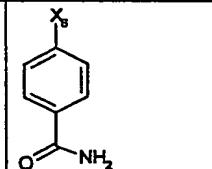
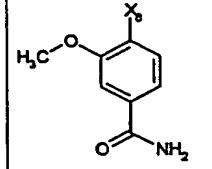
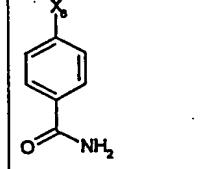
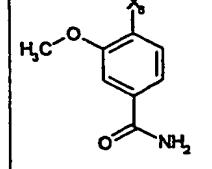
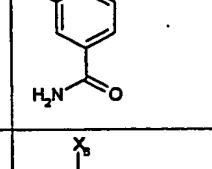
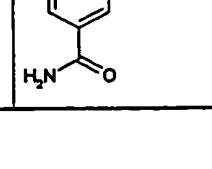
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp.[°C]
227	$\begin{array}{c} X_2 \\ \\ CH_3 \end{array}$	H_3C-X_3	H	R	$\begin{array}{c} CH_3 \\ \\ H_3C-X_3 \\ \\ CH_3 \end{array}$		241
228	$\begin{array}{c} X_2 \\ \\ CH_3 \end{array}$	H_3C-X_3	H	R	$\begin{array}{c} CH_3 \\ \\ H_3C-X_3 \\ \\ CH_3 \end{array}$		
229	$\begin{array}{c} CH_3 \\ \\ X_2 \end{array}$	H_3C-X_3	H	R	$H_3C-CH_2-X_5$		300
230	$\begin{array}{c} CH_3 \\ \\ X_2 \end{array}$	H_3C-X_3	H	R	$H_3C-CH_2-X_5$		200
231	$\begin{array}{c} CH_3 \\ \\ X_2 \end{array}$	$X_3-CH_2-CH_3$	H	R	$H_3C-CH_2-X_5$		232
232	$\begin{array}{c} CH_3 \\ \\ X_2 \end{array}$	$X_3-CH_2-CH_3$	H	R	$H_3C-CH_2-X_5$		149
234	$\begin{array}{c} CH_3 \\ \\ X_2 \end{array}$	H_3C-X_3	H	R			197
235	$\begin{array}{c} CH_3 \\ \\ X_2 \end{array}$	H_3C-X_3	H	R			226

Table 1 (Continued)

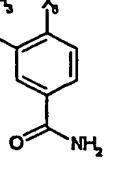
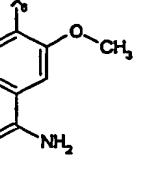
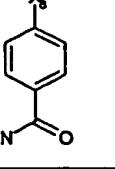
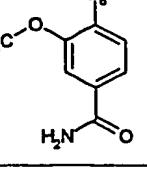
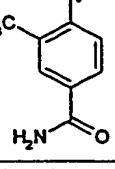
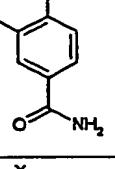
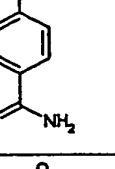
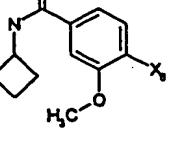
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
236	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	R			182
237	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	R			
238	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$	H	R			
239	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{X}_3 \\ \\ \text{CH}_3 \end{array}$	H	R			
240	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	R			
241	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	R			194
242	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	R			200
243	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	R			156

Table 1 (Continued)

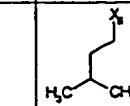
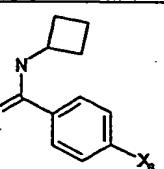
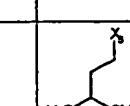
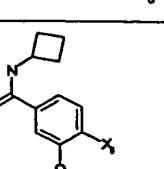
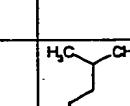
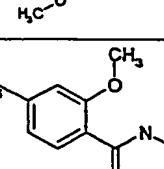
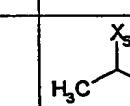
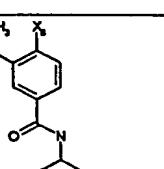
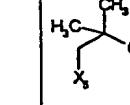
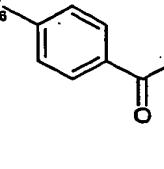
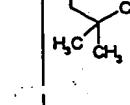
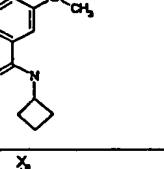
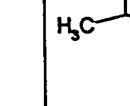
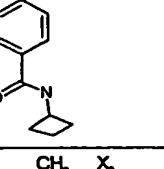
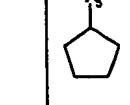
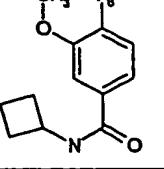
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp.[°C]
244	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{X}_3 - \text{CH}_3$		H	R		 195
245	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{X}_3 - \text{CH}_3$		H	R		 147
246	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$		H	R		
247	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$		H	R		 85
248	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$		H	R		
249	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$		H	R		
250	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$		H	R		 158
251	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$		H	R		 188

Table 1 (Continued)

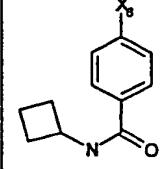
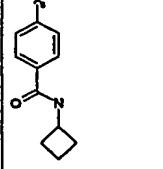
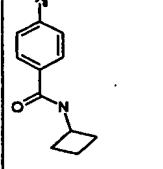
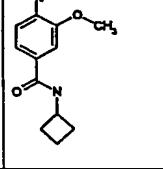
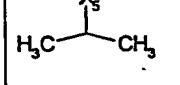
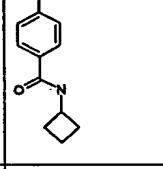
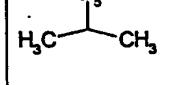
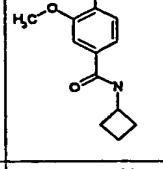
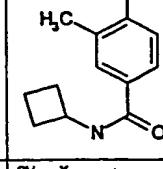
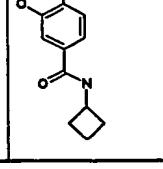
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp.[°C]
252	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	R			245
253	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	R			
254	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	R			128
255	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	R			
256	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{X}_3\text{---CH}_3$	H	R			181
257	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{X}_3\text{---CH}_3$	H	R			217
258	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	R			
259	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\begin{array}{c} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{array}$	H	R			

Table 1 (Continued)

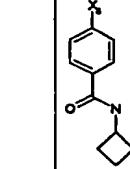
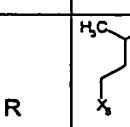
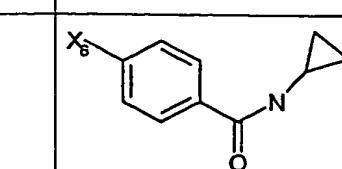
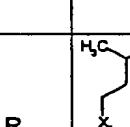
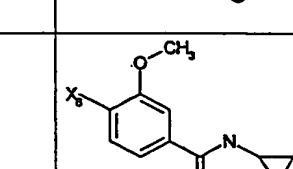
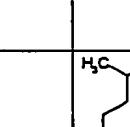
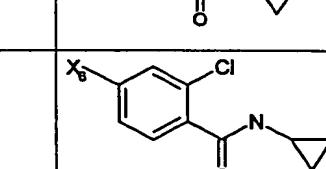
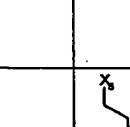
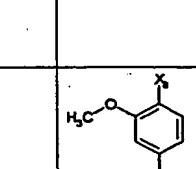
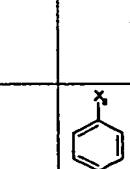
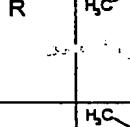
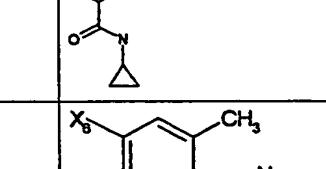
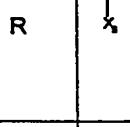
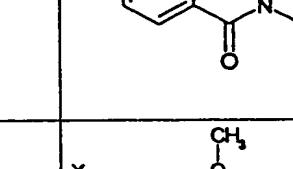
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
260	$\begin{matrix} \text{CH}_3 \\ \\ \text{X}_2 \end{matrix}$	$\text{H}_3\text{C}-\text{X}_3$	H	R			
261	$\begin{matrix} \text{X}_2 \\ \\ \text{CH}_3 \end{matrix}$	$\text{H}_3\text{C}-\text{X}_3$	H	R			230
262	$\begin{matrix} \text{X}_2 \\ \\ \text{CH}_3 \end{matrix}$	$\text{H}_3\text{C}-\text{X}_3$	H	R			193
263	$\begin{matrix} \text{X}_2 \\ \\ \text{CH}_3 \end{matrix}$	$\text{H}_3\text{C}-\text{X}_3$	H	R			
264	$\begin{matrix} \text{CH}_3 \\ \\ \text{X}_2 \end{matrix}$	X_3-CH_3	H	R			152
265	$\begin{matrix} \text{CH}_3 \\ \\ \text{X}_2 \end{matrix}$	X_3-CH_3	H	R			207
266	$\begin{matrix} \text{X}_2 \\ \\ \text{CH}_3 \end{matrix}$	$\text{H}_3\text{C}-\text{X}_3$	H	R			229
267	$\begin{matrix} \text{X}_2 \\ \\ \text{CH}_3 \end{matrix}$	$\text{H}_3\text{C}-\text{X}_3$	H	R			

Table 1 (Continued)

Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp.[°C]
268	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \end{array}$	$\begin{array}{c} \text{CH}_3 \\ \\ \text{O}-\text{C}_6\text{H}_3(\text{X}_8)-\text{C}(=\text{O})-\text{N} \\ \\ \text{C}_6\text{H}_4 \end{array}$	183
269	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	$\begin{array}{c} \text{CH}_3 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \\ \\ \text{X}_8 \end{array}$	$\begin{array}{c} \text{O}-\text{CH}_3 \\ \\ \text{X}_8-\text{C}_6\text{H}_3(\text{X}_8)-\text{C}(=\text{O})-\text{N} \\ \\ \text{C}_6\text{H}_4 \end{array}$	
270	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	$\begin{array}{c} \text{CH}_3 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \\ \\ \text{X}_8 \end{array}$	$\begin{array}{c} \text{X}_8-\text{C}_6\text{H}_3(\text{X}_8)-\text{C}(=\text{O})-\text{N} \\ \\ \text{C}_6\text{H}_4 \end{array}$	161
271	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \end{array}$	$\begin{array}{c} \text{X}_8 \\ \\ \text{C}_6\text{H}_3(\text{X}_8)-\text{C}(=\text{O})-\text{N} \\ \\ \text{C}_6\text{H}_4 \end{array}$	282
272	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	X_3-CH_3	H	R	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \end{array}$	$\begin{array}{c} \text{H}_3\text{C}-\text{O}-\text{C}_6\text{H}_3(\text{X}_8)-\text{C}(=\text{O})-\text{N} \\ \\ \text{C}_6\text{H}_4 \end{array}$	157
273	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	X_3-CH_3	H	R	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \end{array}$	$\begin{array}{c} \text{X}_8 \\ \\ \text{C}_6\text{H}_3(\text{X}_8)-\text{C}(=\text{O})-\text{N} \\ \\ \text{C}_6\text{H}_4 \end{array}$	129
274	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	$\text{X}_5-\text{C}_5\text{H}_9$	$\begin{array}{c} \text{CH}_3 \\ \\ \text{O}-\text{C}_6\text{H}_3(\text{X}_8)-\text{C}(=\text{O})-\text{N} \\ \\ \text{C}_6\text{H}_4 \\ \\ \text{C}_3\text{H}_5 \end{array}$	164
275	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	$\text{X}_5-\text{C}_5\text{H}_9$	$\begin{array}{c} \text{X}_8 \\ \\ \text{C}_6\text{H}_3(\text{X}_8)-\text{C}(=\text{O})-\text{N} \\ \\ \text{C}_6\text{H}_4 \\ \\ \text{C}_3\text{H}_5 \end{array}$	219

Table 1 (Continued)

Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
276			H	R			
277			H	R			200
278			H	R			200
279			H	R			
280			H	R			
281			H	R			
282			H	R			
283			H	R			277

Table 1 (Continued)

Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
284	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	X_5 cyclohexyl	$\begin{array}{c} \text{CH}_3 \\ \\ \text{O} \\ \\ \text{X}_6 \\ \text{C}_6\text{H}_4 \\ \\ \text{O}=\text{C} \\ \text{N} \\ \text{C}_3\text{H}_5 \end{array}$	197
285	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	X_5 cyclohexyl	$\begin{array}{c} \text{X}_6 \\ \text{C}_6\text{H}_4 \\ \\ \text{O}=\text{C} \\ \text{N} \\ \text{C}_3\text{H}_5 \end{array}$	
286	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C}-\text{CH}_2 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \end{array}$	$\begin{array}{c} \text{H}_3\text{C}-\text{N} \\ \\ \text{C}_6\text{H}_4 \\ \\ \text{H}_3\text{C}-\text{O} \\ \text{X}_6 \end{array}$	182
287	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	$\begin{array}{c} \text{H}_3\text{C}-\text{CH}_2 \\ \\ \text{X}_5 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \end{array}$	$\begin{array}{c} \text{O}-\text{CH}_3 \\ \text{X}_6-\text{C}_6\text{H}_4 \\ \\ \text{O}=\text{C} \\ \text{N}-\text{CH}_3 \end{array}$	
288	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	X_3-CH_3	H	R	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C}-\text{CH}_2 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \end{array}$	$\begin{array}{c} \text{CH}_3 \\ \\ \text{O} \\ \\ \text{X}_6-\text{C}_6\text{H}_4 \\ \\ \text{O}=\text{C} \\ \text{N}-\text{CH}_3 \end{array}$	163
289	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	X_3-CH_3	H	R	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C}-\text{CH}_2 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \end{array}$	$\begin{array}{c} \text{X}_6-\text{C}_6\text{H}_4 \\ \\ \text{O}=\text{C} \\ \text{N}-\text{CH}_3 \end{array}$	212
290	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	$\begin{array}{c} \text{H}_3\text{C}-\text{CH}_2 \\ \\ \text{X}_5 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \end{array}$	$\begin{array}{c} \text{CH}_3 \\ \\ \text{O} \\ \\ \text{X}_6-\text{C}_6\text{H}_4 \\ \\ \text{O}=\text{C} \\ \text{N}-\text{CH}_3 \end{array}$	
291	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R	$\begin{array}{c} \text{X}_5 \\ \\ \text{H}_3\text{C}-\text{CH}_2 \\ \\ \text{H}_3\text{C}-\text{CH}_3 \end{array}$	$\begin{array}{c} \text{X}_6-\text{C}_6\text{H}_4 \\ \\ \text{O}-\text{CH}_3 \\ \\ \text{O}=\text{C} \\ \text{N}-\text{CH}_3 \end{array}$	

Table 1 (Continued)

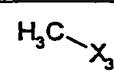
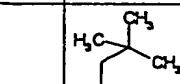
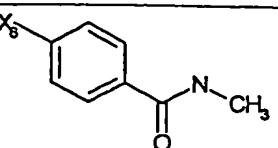
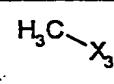
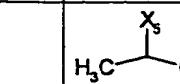
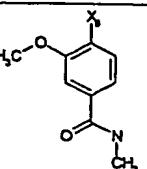
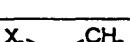
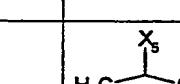
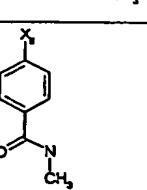
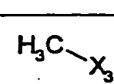
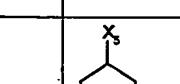
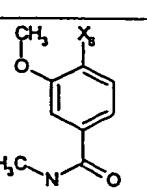
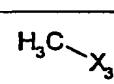
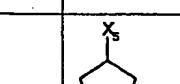
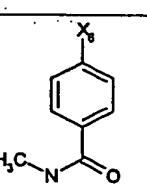
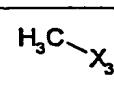
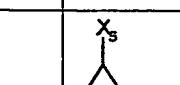
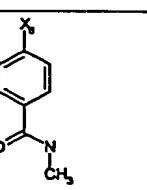
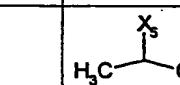
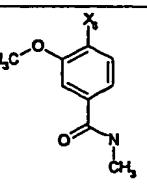
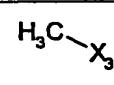
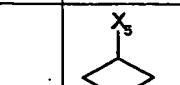
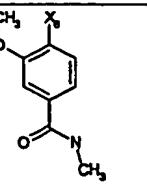
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
292			H	R			
293			H	R			
294			H	R			200
295			H	R			144
296			H	R			221
297			H	R			150
298			H	R			
299			H	R			163

Table 1 (Continued)

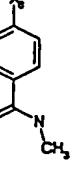
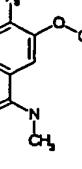
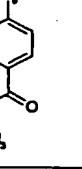
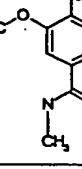
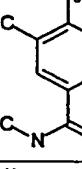
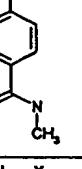
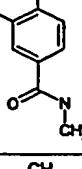
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301	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R			98
302	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	X_3-CH_3	H	R			
303	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	X_3-CH_3	H	R			
304	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R			
305	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R			
306	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R			
307	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C}-\text{X}_3$	H	R			179

Table 1 (Continued)

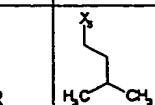
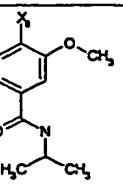
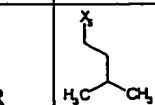
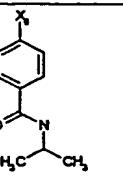
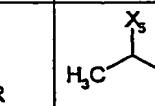
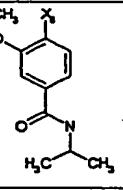
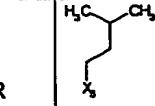
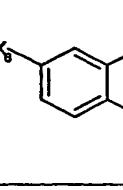
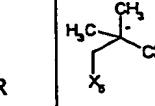
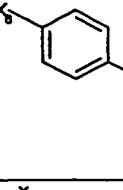
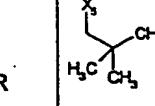
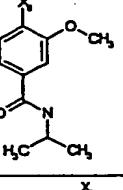
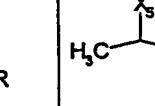
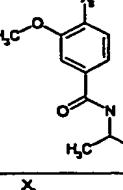
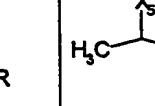
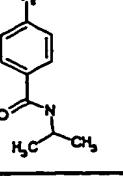
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309	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{X}_3 - \text{CH}_3$		H	R		 231
310	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$		H	R		
311	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$		H	R		 69
312	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$		H	R		
313	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$		H	R		 200
314	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{X}_3 - \text{CH}_3$		H	R		 200
315	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$		H	R		

Table 1 (Continued)

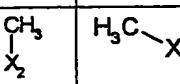
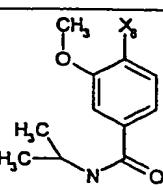
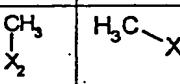
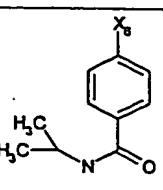
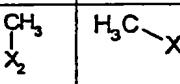
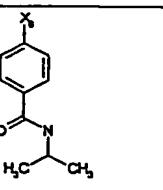
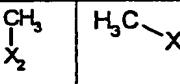
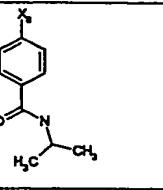
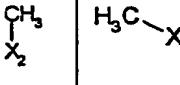
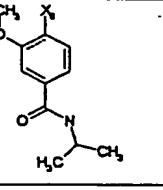
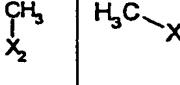
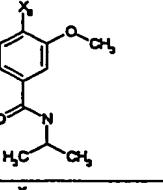
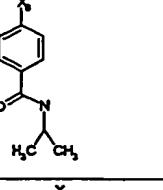
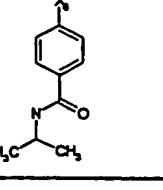
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317			H	R			131
318			H	R			
319			H	R			145
320			H	R			
321			H	R			
322			H	R			149
323			H	R			

Table 1 (Continued)

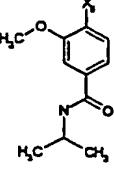
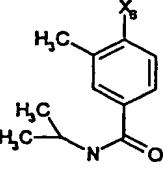
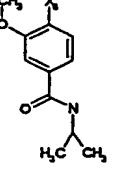
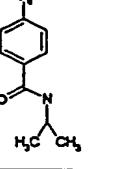
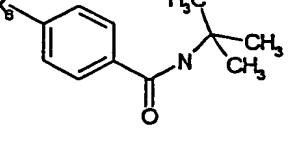
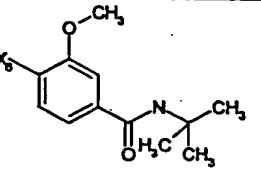
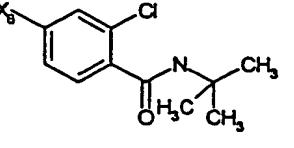
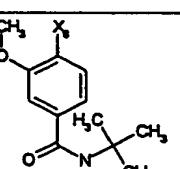
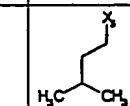
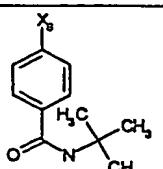
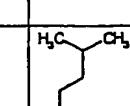
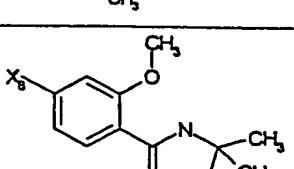
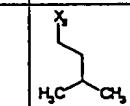
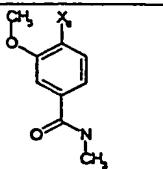
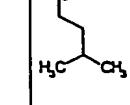
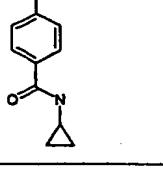
Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
324	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{X}_3 - \text{CH}_3$	H	R			
325	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$	H	R			
326	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$	H	R			
327	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$	H	R			
328	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$	H	R			
329	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$	H	R			176
330	$\begin{array}{c} \text{X}_2 \\ \\ \text{CH}_3 \end{array}$	$\text{H}_3\text{C} - \text{X}_3$	H	R			
331	$\begin{array}{c} \text{CH}_3 \\ \\ \text{X}_2 \end{array}$	$\text{X}_3 - \text{CH}_3$	H	R			

Table 1 (Continued)

Ex.	R ²	R ³	R ⁴	config. R ³ or R ⁴	R ⁵	R ⁶	mp. [°C]
332	$\begin{matrix} \text{CH}_3 \\ \\ \text{X}_2 \end{matrix}$	$\begin{matrix} \text{X}_3 \\ \\ \text{CH}_3 \end{matrix}$		H	R		
333	$\begin{matrix} \text{X}_2 \\ \\ \text{CH}_3 \end{matrix}$	$\begin{matrix} \text{H}_3\text{C} \\ \\ \text{X}_3 \end{matrix}$		H	R		
334	$\begin{matrix} \text{CH}_3 \\ \\ \text{X}_2 \end{matrix}$	$\begin{matrix} \text{X}_3-\text{CH}_3 \\ \\ \text{X}_4-\text{CH}_3 \end{matrix}$					
335	$\begin{matrix} \text{CH}_3 \\ \\ \text{X}_2 \end{matrix}$	$\begin{matrix} \text{X}_5-\text{CH}_3 \\ \\ \text{X}_6-\text{CH}_3 \end{matrix}$					

In the preceding Table the abbreviations X1 to X6 in the groups specified denote the bond which links the particular group to the corresponding group R1 to R6.

As has been found, the compounds of general formula (I) are characterised by their wide range of applications in the therapeutic field. Particular mention should be made of those applications in which the inhibition of specific cell cycle kinases, particularly the inhibiting effect on the proliferation of cultivated human tumour cells 5 but also the proliferation of other cells, such as endothelial cells, for example, plays a part.

As could be demonstrated by FACS analysis, the inhibition of proliferation brought about by the compounds according to the invention is mediated by the arrest of the 10 cells, particularly at the G2/M phase of the cell cycle. The cells arrest, independently of the cells used, for a specific length of time in this phase of the cell cycle before programmed cell death is initiated. An arrest in the G2/M phase of the cell cycle is triggered, for example, by the inhibition of specific cell cycle kinases. Studies in 15 model organisms such as *Schizosaccharomyces pombe* or *Xenopus*, or investigations in human cells have shown that the transition from the G2 phase to mitosis is regulated by the CDK1/cyclin B kinase (Nurse, 1990). This kinase, which is also known as the "mitosis promoting factor" (MPF), phosphorylates and thereby 20 regulates a number of proteins, such as e.g. nuclear lamins, kinesin-like motor proteins, condensins and Golgi matrix proteins, which play an important part in the breakdown of the nuclear envelope, in centrosome separation, the formation of the mitotic spindle apparatus, chromosome condensation and the breakdown of the Golgi apparatus (Nigg. E., 2001). A murine cell line with a temperature-sensitive 25 CDK1 kinase mutant shows a rapid breakdown of the CDK1 kinase and a subsequent arrest in the G2/M phase after a temperature increase (Th'ng et al., 1990). The treatment of human tumour cells with inhibitors against CDK1/cyclin B such as e.g. butyrolactone also leads to an arrest in the G2/M phase and subsequent 30 apoptosis (Nishio, et al. 1996). Another kinase which is involved in the G2 and mitosis phase is polo-like kinase 1 (Plk1), which is responsible for the maturation of the centrosomes, for the activation of the phosphatase Cdc25C, as well as for the activation of the anaphase promoting complex (Glover et al., 1998, Qian, et al., 2001). The injection of Plk1 antibodies leads to a G2 arrest in untransformed cells whereas tumour cells arrest in the mitosis phase (Lane and Nigg, 1996). In addition, the protein kinase aurora B has been described as having an essential function during entry into mitosis. Aurora B phosphorylates histone H3 at Ser11 and thereby

initiates chromosome condensation (Hsu, J.Y. et al., 2000). A specific cell cycle arrest in the G2/M phase may, however, also be triggered e.g. by the inhibition of specific phosphatases such as e.g. Cdc25C (Russell and Nurse, 1986). Yeasts with a defective cdc25 gene arrest in the G2 phase, while overexpression of cdc25 leads 5 to early entry into the mitosis phase (Russell and Nurse, 1987). However, an arrest in the G2/M phase can also be triggered by the inhibition of certain motor proteins, so-capped kinesins such as e.g. Eg5 (Mayer et al., 1999), or by agents which stabilise or destabilise microtubules (e.g. colchicin, taxol, etoposide, vinblastin, vincristin) (Schiff and Horwitz, 1980).

10

In view of their biological properties the compounds of general formula I according to the invention, their isomers and their physiologically acceptable salts are suitable for the treatment of diseases characterised by excessive or abnormal cell proliferation.

15

Such diseases include, for example: viral infections (e.g. HIV and Kaposi's sarcoma); inflammatory and autoimmune diseases (e.g. colitis, arthritis, Alzheimer's disease, glomerulonephritis and wound healing); bacterial, fungal and/or parasitic infections; leukaemias, lymphoma and solid tumours; skin diseases (e.g. psoriasis); bone diseases; cardiovascular diseases (e.g. restenosis and hypertrophy). They are also 20 suitable for protecting proliferating cells (e.g. hair, intestinal, blood and progenitor cells) from damage to their DNA caused by radiation, UV treatment and/or cytostatic treatment (Davis et al., 2001).

25

The new compounds may be used for the prevention, short-term or long-term treatment of the abovementioned diseases, also in combination with other active substances used for the same indications, e.g. cytostatics.

30

The activity of the compounds according to the invention was determined in the cytotoxicity test on cultivated human tumour cells and/or in a FACS analysis, for example on HeLaS3 cells. In both test methods, the compounds exhibited a good to very good activity, i.e. for example an EC₅₀ value in the HeLaS3 cytotoxicity test of less than 5 µmol, generally less than 1 µmol.

Measurement of cytotoxicity on cultivated human tumour cells

To measure the cytotoxicity on cultivated human tumour cells, cells of the cervical 35 cancer tumour cell line HeLaS3 (obtained from American Type Culture Collection

(ATCC)) in Ham's F12 Medium (Life Technologies) and 10% foetal calf serum (Life Technologies) were cultivated and harvested in the logarithmic growth phase. Then the HeLaS3 cells were placed in 96-well plates (Costar) at a density of 1000 cells per well and incubated overnight in an incubator (at 37°C and 5 % CO₂), while on each 5 plate 6 wells were filled only with medium (3 wells as a control of the medium, 3 wells for incubation with reduced AlamarBlue). The active substances were added to the cells in various concentrations (dissolved in DMSO; final concentration: 1%) (in each case as a triple measurement). After 72 hours' incubation, 20 µl of AlamarBlue (AccuMed International) were added to each well, and the cells were incubated for a 10 further 7 hours. As a control, 20 µl of reduced Alamar Blue (AlamarBlue reagent which had been autoclaved for 30 min) were added to 3 wells. After 7 h incubation the colour change of the AlamarBlue reagent in the individual wells was determined in a Perkin Elmer fluorescence spectrophotometer (excitation 530 nm, emission 590 nm, slits 15, integrate time 0.1). The amount of AlamarBlue reagent reacted 15 represents the metabolic activity of the cells. The relative cell activity was calculated as a percentage of the control (HeLa S3 cells without inhibitor) and the active substance concentration which inhibits the cell activity by 50% (IC⁵⁰) was obtained. The values were calculated from the average of three individual measurements, correcting for the control value (medium control).

20

FACS Analysis

Propidium iodide (PI) binds stoichiometrically to double-stranded DNA, and is thus suitable for determining the percentage of cells in the G1, S and G2/M phase of the cell cycle on the basis of the cell DNA content. Cells in the G0 and G1 phase have a 25 diploid DNA content (2N), whereas cells in G2 or mitosis have a 4N DNA content. For PI staining, 0.4 million HeLaS3 cells were seeded, for example, on a 75 cm² cell culture flask, and after 24 h either 1 % DMSO was added as control or the substance was added in various concentrations (in 1% DMSO). The cells were incubated for 24 h with the substance or with DMSO, before the cells were washed with 2 x PBS and 30 detached with trypsin /EDTA. The cells were centrifuged (1000 rpm, 5 min, 4°C), and the cell pellet was washed 2 x with PBS, before the cells were resuspended in 0.1 ml of PBS. Then the cells were fixed with 80% ethanol for 16 hours at 4°C or alternatively for 2 hours at -20°C. The fixed cells (10⁶ cells) were centrifuged (1000 rpm, 5 min, 4°C), washed with PBS and then centrifuged again. The cell pellet was

resuspended in 2 ml of Triton X-100 in 0.25 % PBS, and incubated for 5 min on ice, before 5 ml of PBS were added and the mixture was centrifuged again. The cell pellet was resuspended in 350 μ l of PI stain solution (0.1 mg/ml of Raze A, 10 μ g/ml of presidium iodide in 1 x PBS). The cells were incubated for 20 min in the dark with 5 the stain buffer before being transferred into sample measuring vessels for the FACS scan. The DNA measurement was carried out in a Becton Dickinson FACS Analyzer, with an argon laser (500 mW, emission 488 nm), and the DNA Cell Quest Program (BD). The logarithmic PI fluorescence was determined with a band-pass filter (BP 585/42). The cell populations in the individual phases of the cell cycle were quantified 10 with the ModFit LT program of Becton Dickinson.

The compounds of general formula (I) may be used on their own or combined with other active substances according to the invention, optionally also in conjunction with 15 other pharmacologically active substances. Suitable preparations include for example tablets, capsules, suppositories, solutions, particularly solutions for injection (s.c., i.v., i.m.) and infusion, syrups, emulsions or dispersible powders. The amount of pharmaceutically active compound in each case should be in the range from 0.1 - 90 20 wt.%, preferably 0.5 - 50 wt.% of the total composition, i.e. in amounts which are sufficient to achieve the dosage range given below. The doses specified may, if necessary, be given several times a day.

Suitable tablets may be obtained, for example, by mixing the active substance(s) with known excipients, for example inert diluents such as calcium carbonate, calcium 25 phosphate or lactose, disintegrants such as corn starch or alginic acid, binders such as starch or gelatine, lubricants such as magnesium stearate or talc and/or agents for delaying release, such as carboxymethyl cellulose, cellulose acetate phthalate, or polyvinyl acetate. The tablets may also comprise several layers.

30 Coated tablets may be prepared accordingly by coating cores produced analogously to the tablets with substances normally used for tablet coatings, for example collidone or shellac, gum arabic, talc, titanium dioxide or sugar. To achieve delayed release or prevent incompatibilities the core may also consist of a number of layers. Similarly the tablet coating may consist of a number of layers to achieve delayed 35 release, possibly using the excipients mentioned above for the tablets.

Syrups or elixirs containing the active substances or combinations thereof according to the invention may additionally contain a sweetener such as saccharin, cyclamate, glycerol or sugar and a flavour enhancer, e.g. a flavouring such as vanillin or orange extract. They may also contain suspension adjuvants or thickeners such as sodium carboxymethyl cellulose, wetting agents such as, for example, condensation products of fatty alcohols with ethylene oxide, or preservatives such as p-hydroxybenzoates.

Solutions for injection and infusion are prepared in the usual way, e.g. with the addition of preservatives such as p-hydroxybenzoates, or stabilisers such as alkali metal salts of ethylenediamine tetraacetic acid, optionally using emulsifiers and/or dispersants, while if water is used as the diluent organic solvents may optionally be used as solubilisers or auxiliary solvents, and transferred into injection vials or ampoules or infusion bottles.

15 Capsules containing one or more active substances or combinations of active substances may for example be prepared by mixing the active substances with inert carriers such as lactose or sorbitol and packing them into gelatine capsules.

20 Suitable suppositories may be made for example by mixing with carriers provided for this purpose, such as neutral fats or polyethyleneglycol or the derivatives thereof.

Suitable excipients may be, for example, water, pharmaceutically acceptable organic solvents, such as paraffins (e.g. petroleum fractions), oils of vegetable origin (e.g. groundnut or sesame oil), mono- or polyfunctional alcohols (e.g. ethanol or glycerol), carriers such as e.g. natural mineral powders (e.g. kaolin, clays, talc, chalk), synthetic mineral powders (e.g. highly dispersed silica and silicates), sugar (e.g. glucose, lactose and dextrose), emulsifiers (e.g. lignin, spent sulphite liquors, methylcellulose, starch and polyvinylpyrrolidone) and lubricants (e.g. magnesium stearate, talc, stearic acid and sodium lauryl sulphate).

30 The preparations are administered in the usual way, preferably by oral or transdermal route, particularly preferably by oral route. When administered orally the tablets may, of course, contain additives, such as e.g. sodium citrate, calcium carbonate and dicalcium phosphate together with various additives, such as starch,

preferably potato starch, gelatine and the like, in addition to the abovementioned carriers. Lubricants such as magnesium stearate, sodium laurylsulphate and talc may also be used to form tablets. In the case of aqueous suspensions the active substances may be combined with various flavour enhancers or colourings in

5 addition to the abovementioned excipients.

For parenteral use, solutions of the active substances may be prepared using suitable liquid carrier materials.

The dosage for intravenous use is 1 - 1000 mg per hour, preferably between 5 - 500 mg per hour.

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However, it may optionally be necessary to deviate from the amounts specified, depending on the body weight or method of administration, the individual response to the medication, the nature of the formulation used and the time or interval over which it is administered. Thus, in some cases, it may be sufficient to use less than the 15 minimum quantity specified above, while in other cases the upper limit specified will have to be exceeded. When large amounts are administered it may be advisable to spread them over the day in a number of single doses.

The formulation examples that follow illustrate the present invention without

20 restricting its scope:

Examples of pharmaceutical formulations

A) Tablets per tablet

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active substance	100 mg
lactose	140 mg
corn starch	240 mg
polyvinylpyrrolidone	15 mg
magnesium stearate	5 mg

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500 mg

The finely ground active substance, lactose and some of the corn starch are mixed together. The mixture is screened, then moistened with a solution of polyvinylpyrrolidone in water, kneaded, wet-granulated and dried. The granules, the

remaining corn starch and the magnesium stearate are screened and mixed together. The mixture is compressed to produce tablets of suitable shape and size.

B) Tablets per tablet

5	active substance	80 mg
	lactose	55 mg
	corn starch	190 mg
	microcrystalline cellulose	35 mg
10	polyvinylpyrrolidone	15 mg
	sodium-carboxymethyl starch	23 mg
	magnesium stearate	2 mg
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	400 mg	

15 The finely ground active substance, some of the corn starch, lactose, microcrystalline cellulose and polyvinylpyrrolidone are mixed together, the mixture is screened and worked with the remaining corn starch and water to form a granulate which is dried and screened. The sodiumcarboxymethyl starch and the magnesium stearate are 20 added and mixed in and the mixture is compressed to form tablets of a suitable size.

C) Ampoule solution

25	active substance	50 mg
	sodium chloride	50 mg
	water for inj.	5 ml

30 The active substance is dissolved in water at its own pH or optionally at pH 5.5 to 6.5 and sodium chloride is added to make it isotonic. The solution obtained is filtered free from pyrogens and the filtrate is transferred under aseptic conditions into ampoules which are then sterilised and sealed by fusion. The ampoules contain 5 mg, 25 mg and 50 mg of active substance.